

(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 106 690 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
13.06.2001 Bulletin 2001/24

(51) Int Cl.7: C12N 15/12, C07K 14/47,
C12Q 1/68, C07K 16/18,
G01N 33/53, C07K 19/00,
A61K 39/395, A61K 38/17,
G01N 33/574, G01N 33/68

(21) Application number: 00310408.0

(22) Date of filing: 23.11.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR

Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 23.11.1999 US 447399

(71) Applicant: DIAGNOSTIC PRODUCTS
CORPORATION
Los Angeles California 90045 (US)

(72) Inventors:
• El Shami, A. Said
Camarillo, California 93032 (US)

- Menon, Surendra Nath
Culver City, California 90232 (US)
- French, Cynthia K.
Irvine, California 92612 (US)

(74) Representative:
Campbell, Patrick John Henry et al
J.A. Kemp & Co.,
14 South Square,
Gray's Inn
London WC1R 5LX (GB)

(54) Polynucleotide encoding autoantigens associated with endometriosis

(57) This invention provides a polynucleotide encoding Repro-EN-1.0 and IB1, polypeptides associated with endometriosis. Auto-antibodies against Repro-EN-

1.0 and IB1 have been found in subjects diagnosed with endometriosis. This invention also provides methods of using this polynucleotide and polypeptide.

Description**FIELD OF THE INVENTION**

- 5 [0001] This invention is directed to the field of molecular biology in general, and, more specifically, to a polypeptide associated with endometriosis, an isolated polynucleotide encoding the polypeptide, and methods of using these molecules.

BACKGROUND OF THE INVENTION

- 10 [0002] Endometriosis is a painful disorder that is characterized by the ectopic implantation of functioning endometrial tissue into the abdominal wall and the outer surface of various organs including, most commonly, the lower bowel, ovaries and fallopian tubes. P. Vigano et al. (1991) *Fertility and Sterility* 56:894. Currently, endometriosis-specific genes have not been identified and the events relating to the development of endometriosis are poorly understood. However, 15 several reports suggest that retrograde menstruation linked with abnormal immune function may play a role in establishing ectopic endometrium lesions. T. Ishimaru and H. Masuzaki (1991) *Am. J. Obstet. Gynecol.* 165:210-214. Many attempts to isolate antigens from ectopic endometrium lesions have failed, due to the necrotic nature of the lesions.
- [0003] Endometriosis also is recognized as having an autoimmune component. IgG and IgA auto-antibodies that react with multiple endometrial antigens have been documented in patients with endometriosis. However, attempts to 20 develop IgG or IgA-based assays for the diagnosis of endometriosis has fallen short of fruition. S. Fernandez-Shaw et al., (1996) *Hum. Reprod.* 11:180-1184. R.A. Wild et al. (1991) *Obstetrics and Gynecology* 77:927. Studies have shown that circulating IgG antibodies that bind multiple endometrial proteins can be detected in women with endometriosis to varying degrees. Thirty-five percent to 74% of patients have sera reactive with endometrial proteins. O. Odukoya et al. (1996) *Acta Obstet. Gynecol. Scand.* 75:927-931; J.G. Kim et al. (1995) *Am. J Reprod. Immunol.* 34:80-87; 25 O.A. Odukoya et al. (1995) *Hum. Reprod.* 10:1214-1219. It has also been shown that endometrial antibody titers in patients that respond well to danazol are significantly lower (7/18 (39%) treated patients had elevated titers) than those patients with untreated endometriosis or patients that responded poorly to treatment (17/23 (74%) untreated patients had elevated titers). A. El-Roeiy et al. (1988) *Fertility and Sterility* 50:864-871; H.J. Chihal et al. (1986) *Fertility and Sterility* 46:408-411. In addition, it has been recently reported that women with endometriosis have elevated levels of 30 IL-4, a Th2 mediating cytokine, and that treatment with danazol reduces the levels of IL-4 in women that respond well to treatment. C.-C. Hsu et al. (1997) *Fertility and Sterility* 67:1059-1064.

SUMMARY OF THE INVENTION

- 35 [0004] This invention provides an isolated cDNA molecule and an alternately spliced variant encoding autoantigens associated with endometriosis. The autoantigen is called Repro-EN-1.0. The alternately spliced variant is called IB1. Subjects diagnosed with endometriosis have been found to have antibodies that specifically bind to Repro-EN-1.0 polypeptide and/or a IB1 polypeptide. These antibodies represent a highly sensitive and specific diagnostic marker for endometriosis. Recombinant Repro-EN-1.0 protein and recombinant IB1 protein are useful to detect such antibodies in immunoassays.
- [0005] In one aspect this invention provides a recombinant polynucleotide comprising a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0 (SEQ ID NO:2), or IB1 (SEQ ID NO:4) wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis. In one embodiment the nucleotide sequence is selected from the Repro-EN-1.0 sequence of SEQ ID NO:1 or IB1 sequence of SEQ ID NO: 3. 40 In another embodiment the nucleotide sequence is identical to nucleotides 176 to 2755 of SEQ ID NO:1 or nucleotides 176 to 2986 of SEQ ID NO:3. In another embodiment the polynucleotide further comprises an expression control sequence operatively linked to the nucleotide sequence.
- [0006] In another aspect this invention provides a polynucleotide primer pair which amplifies a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0, or IB1 wherein the epitope specifically binds 45 to antibodies from subjects diagnosed with endometriosis. The pair comprises: 1) a 3' primer of at least 7 nucleotides that specifically hybridizes to a 3' end of the nucleotide sequence or downstream from the sequence, and 2) a 5' primer of at least 7 nucleotides that specifically hybridizes to the 3' end of the complement of the nucleotide sequence or downstream from the complement of the sequence.
- [0007] In another aspect this invention provides a recombinant cell comprising a recombinant polynucleotide comprising an expression control sequence operatively linked to a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0 (SEQ ID NO:2), or IB1 (SEQ ID NO:4), wherein the epitope specifically binds 50 to antibodies from subjects diagnosed with endometriosis.
- [0008] In another aspect this invention provides a method for detecting a target polynucleotide comprising a nucle-

otide sequence selected from Repro-EN-1.0 cDNA (SEQ ID NO: 1) or its complement , or IB1 cDNA (SEQ ID NO:3) or its complement in a sample. The method comprises the steps of: (a) contacting the sample with a polynucleotide probe or primer comprising a sequence of at least 7 nucleotides that specifically hybridizes to the nucleotide sequence and (b) detecting whether the probe or primer has specifically hybridized to the target polynucleotide, whereby specific hybridization provides a detection of the target polynucleotide in the sample.

[0009] In another aspect this invention provides a purified, recombinant Repro-EN-1.0 polypeptide whose amino acid sequence is identical to that of SEQ ID NO:2, or an allelic variant of SEQ ID NO:2, or an IB1 polypeptide whose amino acid sequence is identical to that of SEQ ID No:4, or an allelic variant of SEQ ID No:4.

[0010] In another aspect this invention provides a purified polypeptide comprising an epitope of at least 5 amino acids of Repro-EN-1.0 (SEQ ID NO:2), or an epitope of at least 5 amino acids of IB1 (SEQ ID NO: 4), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.

[0011] In another aspect this invention provides a composition consisting essentially of an antibody that specifically binds to Repro-EN-1.0 polypeptide (SEQ ID NO:2), or IB1 polypeptide (SEQ ID NO:4).

[0012] In another aspect this invention provides a method for detecting a Repro-EN-1.0 polypeptide or IB1 polypeptide in a sample. The method comprises the steps of: (a) contacting the sample with a ligand that specifically binds to the Repro-EN-1.0 polypeptide or IB1 polypeptide and (b) detecting specific binding between the ligand and Repro-EN-1.0 polypeptide or IB1 polypeptide. Specific binding provides a detection of Repro-EN-1.0 polypeptide or IB1 polypeptide in the sample. In one embodiment, the ligand is an antibody.

[0013] In another aspect this invention provides a method diagnosing endometriosis in a subject. The method comprises the steps of: (a) detecting a test amount of an antibody that specifically binds to Repro-EN-1.0 polypeptide or IB1 polypeptide in a sample from the subject; and (b) comparing the test amount with a normal range of the antibody in a control sample from a subject who does not suffer from endometriosis. A test amount above the normal range provides a positive indication in the diagnosis of endometriosis. The sample can be a blood product, e.g., serum, peritoneal fluid, menstrual fluid, vaginal secretion or urine. In one embodiment the antibody is an IgE, IgG or IgG₄ immunoglobulin. In another embodiment the step of detecting comprises capturing the antibody from the sample with an immobilized Repro-EN-1.0 or a peptide comprising an epitope of Repro-EN-1.0 or IB1 or a peptide comprising an epitope of IB1, and detecting captured antibody. The step of detecting captured antibody can comprise contacting the captured antibody with a detectable antibody that specifically binds immunoglobulins and detecting binding between the captured antibody and the detectable antibody. In another embodiment the step of detecting can comprise capturing the antibody from the sample with an immobilized anti-immunoglobulin antibody and detecting captured antibody. The step of detecting captured antibody can comprise contacting the captured antibody with Repro-EN-1.0 or a polypeptide, or IB1 or a peptide comprising an epitope of IB1, comprising an epitope of Repro-EN-1.0 and detecting binding between the captured antibody and the Repro-EN-1.0 or polypeptide or IB1 or polypeptide.

[0014] In another aspect this invention provides a method for use in following the progress of endometriosis in a subject. The method comprises the steps of: (a) detecting first and second amounts of an antibody that specifically bind Repro-EN-1.0 polypeptide, or IB1 polypeptide, in samples from the subject at a first and a second time, respectively; and (b) comparing the first and second amounts. An increase between the first and second amounts indicates progression of the endometriosis and a decrease between the first and second amounts indicates remission of the endometriosis.

[0015] In another aspect this invention provides an isolated MHC-peptide complex comprising at least a portion of an MHC Class I molecule or an MHC Class II molecule, wherein the portion comprises a binding site that specifically binds a peptide having an amino acid binding motif specific to the molecule, and wherein the portion engages in CD4-mediated or CD8-mediated binding to T cells, and a peptide of at least 8 amino acids in a sequence selected from the amino acid sequence of Repro-EN-1.0 (SEQ ID NO:2) or IB1 (SEQ ID NO:4), wherein the peptide comprises the amino acid binding motif and comprises an epitope that specifically binds to a T cell receptor; wherein the complex specifically binds a T cell having a T cell receptor that specifically binds to the epitope, and wherein specific binding induces anergy in the T cell.

[0016] In another aspect this invention provides a method for treating endometriosis in a subject comprising the step of inhibiting an immune response against Repro-EN-1.0 or IB1 in the subject. The method can comprise administering to the subject an immunosuppressant in an amount effective to inhibit the immune response, administering to the subject an isolated MHC-peptide complex of the invention in an amount effective to inhibit the immune response or administering to the subject an anti-idiotypic antibody that specifically binds to an antigen binding site of an antibody that specifically binds to Repro-EN-1.0 or IB1 in an amount effective to inhibit the immune response.

[0017] In another aspect this invention provides a screening method for determining whether a compound increases or decreases the expression of Repro-EN-1.0 in a cell comprising contacting the cell with the compound and determining whether the production of Repro-EN-1.0 mRNA or polypeptide, or IB1 mRNA or polypeptide, are increased or decreased.

[0018] In another aspect this invention provides a method of detecting a chromosomal translocation of a Repro-EN-

1.0 gene or IB1 gene comprising the steps of: a) hybridizing a labeled polynucleotide probe that specifically hybridizes with the Repro-EN-1.0 nucleotide sequence of SEQ ID NO: 1 or its complement, or IB1 nucleotide sequence of SEQ ID NO:3 or its complement, to a chromosome spread from a cell sample to determine the pattern of hybridization and b) determining whether the pattern of hybridization differs from a normal pattern. A difference in the pattern provides detection of a translocation.

5 [0019] In another aspect this invention provides a method of detecting polymorphic forms of Repro-EN-1.0 or IB1 comprising the steps of: a) determining the identity of a nucleotide or amino acid at a selected position within the sequence of a test Repro-EN-1.0 gene or polypeptide, or IB1 gene or polypeptide; b) determining the identity of the nucleotide or amino acid at the corresponding position of native Repro-EN-1.0 (SEQ ID NO: 1 or 2) gene or polypeptide, or IB1 (SEQ No:3 or 4) gene or polypeptide; and c) comparing the identity from the test gene or polynucleotide with the identity of the native gene or polypeptide, whereby a difference in identity indicates that the test polynucleotide is a polymorphic form of Repro-EN-1.0 or IB1.

BRIEF DESCRIPTION OF THE DRAWINGS

15 [0020] Fig. 1 is a northern blot analysis of Repro-EN-1.0 expression in various tissues.

[0021] Fig. 2 is a northern blot analysis of Repro-EN-1.0 expression comparing various normal v. cancerous tissues.

[0022] Fig. 3 is a northern blot analysis of Repro-EN-1.0 expression in various tissue culture cells.

20 DETAILED DESCRIPTION OF THE INVENTION

I. DEFINITIONS

25 [0023] Unless defined otherwise, all technical and scientific terms used herein have the meaning commonly understood by a person skilled in the art to which this invention belongs. The following references provide one of skill with a general definition of many of the terms used in this invention: Singleton et al., DICTIONARY OF MICROBIOLOGY AND MOLECULAR BIOLOGY (2d ed. 1994); THE CAMBRIDGE DICTIONARY OF SCIENCE AND TECHNOLOGY (Walker ed., 1988); THE GLOSSARY OF GENETICS, 5TH ED., R. Rieger et al. (eds.), Springer Verlag (1991); and Hale & Marham, THE HARPER COLLINS DICTIONARY OF BIOLOGY (1991). As used herein, the following terms have the meanings ascribed to them unless specified otherwise.

30 [0024] "Polynucleotide" refers to a polymer composed of nucleotide units. Polynucleotides include naturally occurring nucleic acids, such as deoxyribonucleic acid ("DNA") and ribonucleic acid ("RNA") as well as nucleic acid analogs. Nucleic acid analogs include those which include non-naturally occurring bases, nucleotides that engage in linkages with other nucleotides other than the naturally occurring phosphodiester bond or which include bases attached through linkages other than phosphodiester bonds. Thus, nucleotide analogs include, for example and without limitation, phosphorothioates, phosphorodithioates, phosphorotriesters, phosphoramidates, boranophosphates, methylphosphonates, chiral-methyl phosphonates, 2-O-methyl ribonucleotides, peptide-nucleic acids (PNAs), and the like. Such polynucleotides can be synthesized, for example, using an automated DNA synthesizer. The term "nucleic acid" typically refers to large polynucleotides. The term "oligonucleotide" typically refers to short polynucleotides, generally no greater than about 50 nucleotides. It will be understood that when a nucleotide sequence is represented by a DNA sequence (i.e., A, T, G, C), this also includes an RNA sequence (i.e., A, U, G, C) in which "U" replaces "T."

35 [0025] "cDNA" refers to a DNA that is complementary or identical to an mRNA, in either single stranded or double stranded form.

40 [0026] Conventional notation is used herein to describe polynucleotide sequences: the left-hand end of a single-stranded polynucleotide sequence is the 5'-end; the left-hand direction of a double-stranded polynucleotide sequence is referred to as the 5'-direction. The direction of 5' to 3' addition of nucleotides to nascent RNA transcripts is referred to as the transcription direction. The DNA strand having the same sequence as an mRNA is referred to as the "coding strand"; sequences on the DNA strand having the same sequence as an mRNA transcribed from that DNA and which are located 5' to the 5'-end of the RNA transcript are referred to as "upstream sequences"; sequences on the DNA strand having the same sequence as the RNA and which are 3' to the 3' end of the coding RNA transcript are referred to as "downstream sequences."

45 [0027] "Complementary" refers to the topological compatibility or matching together of interacting surfaces of two polynucleotides. Thus, the two molecules can be described as complementary, and furthermore, the contact surface characteristics are complementary to each other. A first polynucleotide is complementary to a second polynucleotide if the nucleotide sequence of the first polynucleotide is identical to the nucleotide sequence of the polynucleotide binding partner of the second polynucleotide. Thus, the polynucleotide whose sequence 5'-TATACT-3' is complementary to a polynucleotide whose sequence is 5'-GTATA-3'.

50 [0028] A nucleotide sequence is "substantially complementary" to a reference nucleotide sequence if the sequence

complementary to the subject nucleotide sequence is substantially identical to the reference nucleotide sequence.

[0029] "Encoding" refers to the inherent property of specific sequences of nucleotides in a polynucleotide, such as a gene, a cDNA, or an mRNA, to serve as templates for synthesis of other polymers and macromolecules in biological processes having either a defined sequence of nucleotides (i.e., rRNA, tRNA and mRNA) or a defined sequence of amino acids and the biological properties resulting therefrom. Thus, a gene encodes a protein if transcription and translation of mRNA produced by that gene produces the protein in a cell or other biological system. Both the coding strand, the nucleotide sequence of which is identical to the mRNA sequence and is usually provided in sequence listings, and non-coding strand, used as the template for transcription, of a gene or cDNA can be referred to as encoding the protein or other product of that gene or cDNA. Unless otherwise specified, a "nucleotide sequence encoding an amino acid sequence" includes all nucleotide sequences that are degenerate versions of each other and that encode the same amino acid sequence. Nucleotide sequences that encode proteins and RNA may include introns.

[0030] "Recombinant polynucleotide" refers to a polynucleotide having sequences that are not naturally joined together. An amplified or assembled recombinant polynucleotide may be included in a suitable vector, and the vector can be used to transform a suitable host cell. A host cell that comprises the recombinant polynucleotide is referred to as a "recombinant host cell." The gene is then expressed in the recombinant host cell to produce, e.g., a "recombinant polypeptide." A recombinant polynucleotide may serve a non-coding function (e.g., promoter, origin of replication, ribosome-binding site, etc.) as well.

[0031] "Expression control sequence" refers to a nucleotide sequence in a polynucleotide that regulates the expression (transcription and/or translation) of a nucleotide sequence operatively linked thereto. "Operatively linked" refers to a functional relationship between two parts in which the activity of one part (e.g., the ability to regulate transcription) results in an action on the other part (e.g., transcription of the sequence). Expression control sequences can include, for example and without limitation, sequences of promoters (e.g., inducible or constitutive), enhancers, transcription terminators, a start codon (i.e., ATG), splicing signals for introns, and stop codons.

[0032] "Expression vector" refers to a vector comprising a recombinant polynucleotide comprising expression control sequences operatively linked to a nucleotide sequence to be expressed. An expression vector comprises sufficient cis-acting elements for expression; other elements for expression can be supplied by the host cell or *in vitro* expression system. Expression vectors include all those known in the art, such as cosmids, plasmids (e.g., naked or contained in liposomes) and viruses that incorporate the recombinant polynucleotide.

[0033] "Amplification" refers to any means by which a polynucleotide sequence is copied and thus expanded into a larger number of polynucleotide molecules, e.g., by reverse transcription, polymerase chain reaction, and ligase chain reaction.

[0034] "Primer" refers to a polynucleotide that is capable of specifically hybridizing to a designated polynucleotide template and providing a point of initiation for synthesis of a complementary polynucleotide. Such synthesis occurs when the polynucleotide primer is placed under conditions in which synthesis is induced, i.e., in the presence of nucleotides, a complementary polynucleotide template, and an agent for polymerization such as DNA polymerase. A primer is typically single-stranded, but may be double-stranded. Primers are typically deoxyribonucleic acids, but a wide variety of synthetic and naturally occurring primers are useful for many applications. A primer is complementary to the template to which it is designed to hybridize to serve as a site for the initiation of synthesis, but need not reflect the exact sequence of the template. In such a case, specific hybridization of the primer to the template depends on the stringency of the hybridization conditions. Primers can be labeled with, e.g., chromogenic, radioactive, or fluorescent moieties and used as detectable moieties.

[0035] "Probe," when used in reference to a polynucleotide, refers to a polynucleotide that is capable of specifically hybridizing to a designated sequence of another polynucleotide. A probe specifically hybridizes to a target complementary polynucleotide, but need not reflect the exact complementary sequence of the template. In such a case, specific hybridization of the probe to the target depends on the stringency of the hybridization conditions. Probes can be labeled with, e.g., chromogenic, radioactive, or fluorescent moieties and used as detectable moieties.

[0036] A first sequence is an "antisense sequence" with respect to a second sequence if a polynucleotide whose sequence is the first sequence specifically hybridizes with a polynucleotide whose sequence is the second sequence.

[0037] "Hybridizing specifically to" or "specific hybridization" or "selectively hybridize to", refers to the binding, duplexing, or hybridizing of a nucleic acid molecule preferentially to a particular nucleotide sequence under stringent conditions when that sequence is present in a complex mixture (e.g., total cellular) DNA or RNA.

[0038] The term "stringent conditions" refers to conditions under which a probe will hybridize preferentially to its target subsequence, and to a lesser extent to, or not at all to, other sequences. "Stringent hybridization" and "stringent hybridization wash conditions" in the context of nucleic acid hybridization experiments such as Southern and northern hybridizations are sequence dependent, and are different under different environmental parameters. An extensive guide to the hybridization of nucleic acids is found in Tijssen (1993) *Laboratory Techniques in Biochemistry and Molecular Biology--Hybridization with Nucleic Acid Probes* part I chapter 2 "Overview of principles of hybridization and the strategy of nucleic acid probe assays", Elsevier, New York. Generally, highly stringent hybridization and wash

conditions are selected to be about 5° C lower than the thermal melting point (Tm) for the specific sequence at a defined ionic strength and pH. The Tm is the temperature (under defined ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly matched probe. Very stringent conditions are selected to be equal to the Tm for a particular probe.

- 5 [0039] An example of stringent hybridization conditions for hybridization of complementary nucleic acids which have more than 100 complementary residues on a filter in a Southern or northern blot is 50% formalin with 1 mg of heparin at 42° C, with the hybridization being carried out overnight. An example of highly stringent wash conditions is 0.15 M NaCl at 72° C for about 15 minutes. An example of stringent wash conditions is a 0.2X SSC wash at 65° C for 15 minutes (see, Sambrook et al. for a description of SSC buffer). Often, a high stringency wash is preceded by a low stringency wash to remove background probe signal. An example medium stringency wash for a duplex of, e.g., more than 100 nucleotides, is 1x SSC at 45° C for 15 minutes. An example low stringency wash for a duplex of, e.g., more than 100 nucleotides, is 4-6x SSC at 40° C for 15 minutes. In general, a signal to noise ratio of 2x (or higher) than that observed for an unrelated probe in the particular hybridization assay indicates detection of a specific hybridization.

- 10 [0040] "Polypeptide" refers to a polymer composed of amino acid residues, related naturally occurring structural variants, and synthetic non-naturally occurring analogs thereof linked via peptide bonds, related naturally occurring structural variants, and synthetic non-naturally occurring analogs thereof. Synthetic polypeptides can be synthesized, for example, using an automated polypeptide synthesizer. The term "protein" typically refers to large polypeptides. The term "peptide" typically refers to short polypeptides.

- 15 [0041] Conventional notation is used herein to portray polypeptide sequences: the left-hand end of a polypeptide sequence is the amino-terminus; the right-hand end of a polypeptide sequence is the carboxyl-terminus.

- 20 [0042] "Conservative substitution" refers to the substitution in a polypeptide of an amino acid with a functionally similar amino acid. The following six groups each contain amino acids that are conservative substitutions for one another:

- 25 1) Alanine (A), Serine (S), Threonine (T);
 2) Aspartic acid (D), Glutamic acid (E);
 3) Asparagine (N), Glutamine (Q);
 4) Arginine (R), Lysine (K);
 5) Isoleucine (I), Leucine (L), Methionine (M), Valine (V); and
 30 6) Phenylalanine (F), Tyrosine (Y), Tryptophan (W).

- [0043] "Allelic variant" refers to any of two or more polymorphic forms of a gene occupying the same genetic locus. Allelic variations arise naturally through mutation, and may result in phenotypic polymorphism within populations. Gene mutations can be silent (no change in the encoded polypeptide) or may encode polypeptides having altered amino acid sequences. "Allelic variants" also refer to cDNAs derived from mRNA transcripts of genetic allelic variants, as well as the proteins encoded by them.

- [0044] Terms used to describe sequence relationships between two or more nucleotide sequences or amino acid sequences include "reference sequence," "selected from," "comparison window," "identical," "percentage of sequence identity," "substantially identical," "complementary," and "substantially complementary."

- 40 [0045] The terms "identical" or percent "identity," in the context of two or more polynucleotide or polypeptide sequences, refer to two or more sequences or sub-sequences that are the same or have a specified percentage of nucleotides or amino acid residues that are the same, when compared and aligned for maximum correspondence, as measured using one of the following sequence comparison algorithms or by visual inspection.

- [0046] The phrase "substantially identical," in the context of two nucleic acids or polypeptides, refers to two or more sequences or sub-sequences that have at least 60%, 80%, 90%, 95% or 98% nucleotide or amino acid residue identity, when compared and aligned for maximum correspondence, as measured using one of the following sequence comparison algorithms or by visual inspection. Preferably, the substantial identity exists over a region of the sequences that is at least about 50 residues in length, more preferably over a region of at least about 100 residues, and most preferably the sequences are substantially identical over at least about 150 residues. In a most preferred embodiment, the sequences are substantially identical over the entire length of the coding regions.

- [0047] For sequence comparison, typically one sequence acts as a reference sequence, to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are input into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. The sequence comparison algorithm then calculates the percent sequence identity for the test sequence(s) relative to the reference sequence, based on the designated program parameters.

- [0048] Optimal alignment of sequences for comparison can be conducted, e.g., by the local homology algorithm of Smith & Waterman, *Adv. Appl. Math.* 2:482 (1981), by the homology alignment algorithm of Needleman & Wunsch, *J. Mol. Biol.* 48:443 (1970), by the search for similarity method of Pearson & Lipman, *Proc. Nat'l. Acad. Sci. USA* 85:2444

(1988), by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by visual inspection (see generally Ausubel *et al.*, *supra*).

[0049] One example of a useful algorithm is PILEUP. PILEUP creates a multiple sequence alignment from a group of related sequences using progressive, pairwise alignments to show relationship and percent sequence identity. It also plots a tree or dendrogram showing the clustering relationships used to create the alignment. PILEUP uses a simplification of the progressive alignment method of Feng & Doolittle, *J. Mol. Evol.* 35:351-360 (1987). The method used is similar to the method described by Higgins & Sharp, *CABIOS* 5:151-153 (1989). The program can align up to 300 sequences, each of a maximum length of 5,000 nucleotides or amino acids. The multiple alignment procedure begins with the pairwise alignment of the two most similar sequences, producing a cluster of two aligned sequences. This cluster is then aligned to the next most related sequence or cluster of aligned sequences. Two clusters of sequences are aligned by a simple extension of the pairwise alignment of two individual sequences. The final alignment is achieved by a series of progressive, pairwise alignments. The program is run by designating specific sequences and their amino acid or nucleotide coordinates for regions of sequence comparison and by designating the program parameters. For example, a reference sequence can be compared to other test sequences to determine the percent sequence identity relationship using the following parameters: default gap weight (3.00), default gap length weight (0.10), and weighted end gaps.

[0050] Another example of algorithm that is suitable for determining percent sequence identity and sequence similarity is the BLAST algorithm, which is described in Altschul *et al.*, *J. Mol. Biol.* 215:403-410 (1990). Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (<http://www.ncbi.nlm.nih.gov/>). This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul *et al.*, *supra*). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are then extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a word-length (W) of 11, an expectation (E) of 10, M=5, N=-4, and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a word-length (W) of 3, an expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff & Henikoff, *Proc. Nat'l. Acad. Sci. USA* 89:10915 (1989)).

[0051] In addition to calculating percent sequence identity, the BLAST algorithm also performs a statistical analysis of the similarity between two sequences (see, e.g., Karlin & Altschul, *Proc. Proc. Natl. Acad. Sci. USA* 90:5873-5787 (1993)). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur by chance. For example, a nucleic acid is considered similar to a reference sequence if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.1, more preferably less than about 0.01, and most preferably less than about 0.001.

[0052] A further indication that two nucleic acid sequences or polypeptides are substantially identical is that the polypeptide encoded by the first nucleic acid is immunologically cross reactive with the polypeptide encoded by the second nucleic acid, as described below. Thus, a polypeptide is typically substantially identical to a second polypeptide, for example, where the two peptides differ only by conservative substitutions. Another indication that two nucleic acid sequences are substantially identical is that the two molecules hybridize to each other under stringent conditions, as described herein.

[0053] An "affinity agent" is a compound that specifically or non-specifically binds to a target molecule. Affinity agents that non-specifically bind to a molecule include, for example, anion or cation exchange resins, or materials that bind hydrophobic or hydrophilic molecules, or metal ions.

[0054] A "ligand" is a compound that specifically binds to a target molecule.

[0055] A "receptor" is compound that specifically binds to a ligand.

[0056] "Antibody" refers to a polypeptide ligand substantially encoded by an immunoglobulin gene or immunoglobulin genes, or fragments thereof, which specifically binds and recognizes an epitope (e.g., an antigen). The recognized immunoglobulin genes include the kappa and lambda light chain constant region genes, the alpha, gamma, delta, epsilon and mu heavy chain constant region genes, and the myriad immunoglobulin variable region genes. Antibodies exist, e.g., as intact immunoglobulins or as a number of well characterized fragments produced by digestion with various

peptidases. This includes, e.g., Fab' and F(ab)'2 fragments. The term "antibody," as used herein, also includes antibody fragments either produced by the modification of whole antibodies or those synthesized *de novo* using recombinant DNA methodologies. It also includes polyclonal antibodies, monoclonal antibodies, chimeric antibodies and humanized antibodies. "Fc" portion of an antibody refers to that portion of an immunoglobulin heavy chain that comprises one or

5 more heavy chain constant region domains, CH1, CH2 and CH3, but does not include the heavy chain variable region.

[0057] A ligand or a receptor (e.g., an antibody) "specifically binds to" or "is specifically immunoreactive with" a compound analyte when the ligand or receptor functions in a binding reaction which is determinative of the presence of the analyte in a sample of heterogeneous compounds. Thus, under designated assay (e.g., immunoassay) conditions, the ligand or receptor binds preferentially to a particular analyte and does not bind in a significant amount to other compounds present in the sample. For example, a polynucleotide specifically binds under hybridization conditions to an analyte polynucleotide comprising a complementary sequence; an antibody specifically binds under immunoassay conditions to an antigen analyte bearing an epitope against which the antibody was raised; and an adsorbent specifically binds to an analyte under proper elution conditions.

[0058] "Immunoassay" refers to a method of detecting an analyte in a sample in which specificity for the analyte is conferred by the specific binding between an antibody and a ligand. This includes detecting an antibody analyte through specific binding between the antibody and a ligand. See Harlow and Lane (1988) *Antibodies, A Laboratory Manual*, Cold Spring Harbor Publications, New York, for a description of immunoassay formats and conditions that can be used to determine specific immunoreactivity.

[0059] "Vaccine" refers to an agent or composition containing an agent effective to confer a therapeutic degree of immunity on an organism while causing only very low levels of morbidity or mortality. Methods of making vaccines are, of course, useful in the study of the immune system and in preventing and treating animal or human disease.

[0060] An "immunogenic amount" is an amount effective to elicit an immune response in a subject.

[0061] "Substantially pure" or "isolated" means an object species is the predominant species present (i.e., on a molar basis, more abundant than any other individual macromolecular species in the composition), and a substantially purified fraction is a composition wherein the object species comprises at least about 50% (on a molar basis) of all macromolecular species present. Generally, a substantially pure composition means that about 80% to 90% or more of the macromolecular species present in the composition is the purified species of interest. The object species is purified to essential homogeneity (contaminant species cannot be detected in the composition by conventional detection methods) if the composition consists essentially of a single macromolecular species. Solvent species, small molecules (<500 Daltons), stabilizers (e.g., BSA), and elemental ion species are not considered macromolecular species for purposes of this definition.

[0062] "Naturally-occurring" as applied to an object refers to the fact that the object can be found in nature. For example, a polypeptide or polynucleotide sequence that is present in an organism (including viruses) that can be isolated from a source in nature and which has not been intentionally modified by man in the laboratory is naturally-occurring.

[0063] "Detecting" refers to determining the presence, absence, or amount of an analyte in a sample, and can include quantifying the amount of the analyte in a sample or per cell in a sample.

[0064] "Detectable moiety" or a "label" refers to a composition detectable by spectroscopic, photochemical, biochemical, immunochemical, or chemical means. For example, useful labels include ^{32}P , ^{35}S , fluorescent dyes, electron-dense reagents, enzymes (e.g., as commonly used in an ELISA), biotin-streptavidin, dioxigenin, haptens and proteins for which antisera or monoclonal antibodies are available, or nucleic acid molecules with a sequence complementary to a target. The detectable moiety often generates a measurable signal, such as a radioactive, chromogenic, or fluorescent signal, that can be used to quantitate the amount of bound detectable moiety in a sample. The detectable moiety can be incorporated in or attached to a primer or probe either covalently, or through ionic, van der Waals or hydrogen bonds, e.g., incorporation of radioactive nucleotides, or biotinylated nucleotides that are recognized by streptavidin. The detectable moiety may be directly or indirectly detectable. Indirect detection can involve the binding of a second directly or indirectly detectable moiety to the detectable moiety. For example, the detectable moiety can be the ligand of a binding partner, such as biotin, which is a binding partner for streptavidin, or a nucleotide sequence, which is the binding partner for a complementary sequence, to which it can specifically hybridize. The binding partner may itself be directly detectable, for example, an antibody may be itself labeled with a fluorescent molecule. The binding partner also may be indirectly detectable, for example, a nucleic acid having a complementary nucleotide sequence can be a part of a branched DNA molecule that is in turn detectable through hybridization with other labeled nucleic acid molecules. (See, e.g., PD. Fahrlander and A. Klausner, *Bio/Technology* (1988) 6:1165.) Quantitation of the signal is achieved by, e.g., scintillation counting, densitometry, or flow cytometry.

[0065] "Linker" refers to a molecule that joins two other molecules, either covalently, or through ionic, van der Waals or hydrogen bonds, e.g., a nucleic acid molecule that hybridizes to one complementary sequence at the 5' end and to another complementary sequence at the 3' end, thus joining two non-complementary sequences.

[0066] "Pharmaceutical composition" refers to a composition suitable for pharmaceutical use in a mammal. A phar-

maceutical composition comprises a pharmacologically effective amount of an active agent and a pharmaceutically acceptable carrier. "Pharmacologically effective amount" refers to that amount of an agent effective to produce the intended pharmacological result. "Pharmaceutically acceptable carrier" refers to any of the standard pharmaceutical carriers, buffers, and excipients, such as a phosphate buffered saline solution, 5% aqueous solution of dextrose, and emulsions, such as an oil/water or water/oil emulsion, and various types of wetting agents and/or adjuvants. Suitable pharmaceutical carriers and formulations are described in *Remington's Pharmaceutical Sciences*, 19th Ed. (Mack Publishing Co., Easton, 1995). Preferred pharmaceutical carriers depend upon the intended mode of administration of the active agent. Typical modes of administration include enteral (e.g., oral) or parenteral (e.g., subcutaneous, intramuscular, intravenous or intraperitoneal injection; or topical, transdermal, or transmucosal administration). A "pharmaceutically acceptable salt" is a salt that can be formulated into a compound for pharmaceutical use including, e.g., metal salts (sodium, potassium, magnesium, calcium, etc.) and salts of ammonia or organic amines.

[0067] "Small organic molecule" refers to organic molecules of a size comparable to those organic molecules generally used in pharmaceuticals. The term excludes organic biopolymers (e.g., proteins, nucleic acids, etc.). Preferred small organic molecules range in size up to about 5000 Da, up to about 2000 Da, or up to about 1000 Da.

[0068] A "subject" of diagnosis or treatment is a human or non-human mammal. Non-human mammals subject to diagnosis or treatment include, for example, primates, ungulates, canines and felines.

[0069] "Treatment" refers to prophylactic treatment or therapeutic treatment.

[0070] A "prophylactic" treatment is a treatment administered to a subject who does not exhibit signs of a disease or exhibits only early signs for the purpose of decreasing the risk of developing pathology.

[0071] A "therapeutic" treatment is a treatment administered to a subject who exhibits signs of pathology for the purpose of diminishing or eliminating those signs.

[0072] "Diagnostic" means identifying the presence or nature of a pathologic condition. Diagnostic methods differ in their sensitivity and specificity. The "sensitivity" of a diagnostic assay is the percentage of diseased individuals who test positive (percent of true positives). The "specificity" of a diagnostic assay is 1 minus the false positive rate, where the false positive rate is defined as the proportion of those without the disease who test positive. While a particular diagnostic method may not provide a definitive diagnosis of a condition, it suffices if the method provides a positive indication that aids in diagnosis.

[0073] "Prognostic" means predicting the probable development (e.g., severity) of a pathologic condition.

[0074] "Test amount" refers to an amount of an analyte in a subject sample, which is then compared to a normal amount of the analyte in a sample (e.g., from a healthy individual) such that the relative comparison of the values provides a reference value for diagnosing a designated disease. Depending upon the method of detection, the test amount may be a determination of the amount of the analyte, but it is not necessarily an amount. The test amount may also be a relative value, such as a plus or a minus score, and also includes an amount indicating the presence or absence of the analyte in a sample.

[0075] "Normal amount" refers to an amount or a range of an analyte in a biological sample that indicates health or lack of pathology.

[0076] "Diagnostic amount" refers to an amount of an analyte in a subject sample that is consistent with a particular diagnosis for a designated disease.

[0077] "Prognostic amount" refers to an amount or range of an analyte in a subject sample that is consistent with a particular prognosis for a designated disease.

[0078] "Plurality" means at least two.

[0079] An "epitope" is portion of a molecule that specifically binds to an antibody or a T cell receptor. An peptide epitope generally comprises a sequence of at least 6 amino acids from a polypeptide, although longer and shorter peptides can constitute epitopes.

[0080] "MHC Class I molecule" refers to a heterodimer found on the surface of cells that present processed antigenic peptides to T cells. The molecule comprises an α chain and a β -microglobulin chain. The α chain contains the antigenic peptide binding site in the $\alpha 1$ and $\alpha 2$ domains. The α chain also contains a transmembrane portion that can be removed without eliminating antigen binding.

[0081] "MHC Class II molecule" refers to a heterodimer found on the surface of cells that present processed antigenic peptide to T cells. It comprises an α chain and a β chain. The antigenic peptide binding site is located in the $\alpha 1$ domain of the α chain and the $\beta 1$ domain of the β chain. However, a single α chain or β chain suffices to bind an antigenic peptide. The α chain and β chain also contain transmembrane regions that can be removed without eliminating antigenic peptide binding function.

[0082] "T cell receptor" refers to a heterodimer found on the surface of T cells comprising an α chain and a β chain or a γ and a δ chain. T cell receptors recognize processed antigens associated with MHC molecules.

II. cDNA ENCODING REPRO-EN-1.0 and IB1

[0083] We have isolated a cDNA molecule encoding an autoantigen associated with endometriosis. The autoantigen is called Repro-EN-1.0. The presence of antibodies that specifically bind to an epitope of the Repro-EN-1.0 polypeptide is a highly sensitive and specific diagnostic marker for endometriosis.

[0084] Polynucleotides encoding full-length Repro-EN-1.0 are useful in recombinant production Repro-EN-1.0 or immunogenic fragments of it. Fragments of polynucleotides encoding Repro-EN-1.0 are useful as probes to detect Repro-EN-1.0 mRNA from certain cell types suspected to be cancerous. Fragments also are useful as primers for amplification of sequences from Repro-EN-1.0.

[0085] The Repro-EN-1.0 polypeptide and immunogenic fragments of it are useful as positive controls in diagnostic assays to detect antibodies that specifically bind to Repro-EN-EN-1.0 from patient serum samples. The polypeptides also are useful as immunogens for eliciting production of antibodies against epitopes of the protein.

[0086] The nucleotide sequence (SEQ ID NO:1) and deduced amino acid sequence (SEQ ID NO:2) of Repro-EN-1.0 follow:

15

20

25

30

35

40

45

50

55

1 CGGGCGGGCTTCAGGGGCCAGGCGCCGCTGC TGCCACCGCCATCTAACGCTGCGCCCTG 60
 5 GCCGGGCCGAAGTCCCCGGGTCCGCGGCCACGACGGTGGCGGTAGATTGCACGCCGGAC

61 GAGGCCCGGCGCGCGATGGTGCCGGTGC GGCTCGGGTGTAAACGGGTGTCCCCTCCC 120
 10 CTCCGGGCCGCGGCCATACCACGGCCACGCCAGGCCACAACTTGCCACAGGGGAGGG

121 CCTCCTCCCCCTCCCCACGCGGTGGTCTCCCCCTCCACCCGGCTCAGGCAGAGCCATGTC 180
 15 GGAGGAGGGGAGGGGGTGCGCCACCAGAGGGAGGGTGGGCCAGTCCGTCTCGGTACAG

181 TCAGGGGTGGCTCC TACCCACACCTGTTGGGACGTGAGGAAAAGGTTCTCGGGCTGGA 240
 20 AGCCCCACCGAGGAATGGGTGTGGACAACACCTGCACTCCTTTCCAAGGAGCCGACCT

R G G S Y P H L L V D V R K R F L G L E

25 GGACCCGTCCCGGCTCGGAGTCGCTACCTGGGAAGAAGAGAATTATCAAAGATTAAG 300
 241 CCTGGCAGGGCCGACGCCCTAGCGATGGACCCCTCTCTAAATAGGTTCTAAATTT

D P S R L R S R Y L G R R E F I Q R L K

30 ACTTGAAAGCAACCTTAATGTGCATGATGGTTGTTAACATAATCTGTTGGAATGACAC 360
 301 TGAAACTTCGTTGGATTACACGTACTACCAACACAATTATGTTAGACAACCTTACTGTG

L E A T L N V H D G C V N T I C W N D T

35 TGGAGAAATATTTATCTGGCTCAGATGACACCAATTAGTAATTAGTAATCCTTACAG 420
 361 ACCTCTTATATAAAATAGACCGAGTCTACTGTGGTTAACATTAATCATTAGGAATGTC

G E Y I L S G S D D T K L V I S N P Y S

40 CAGAAAGGTTTGACAACAATTGTTAGGGCACCGAGCAAACATATTAGTCAAAGTT 480
 45 GTCTTTCCAAAATGTTAGCAAGTCCCGTGGCTGTTGTATAATCACGTTCAA

R K V L T T I R S G H R A N I F S A K F

50

481 CTTACCTTGTACAAATGATAAACAGATTGTATCCTGCTCTGGAGATGGAGTAATATTTA + 540
 GAATGGAACATGTTACTATTTGTCACATAGGACGAGACCTCACCTCATTATAAAAT

5 L P C T N D K O I V S C S G D G V I F Y

10 TACCAACGTTGAGCAAGATGCAGAAAACCAACAGACAATGCCAATTACGTGTTCATTATGG + 600
 ATGGTTGCAACTCGTCTACGTCTTGGTTCTGTACGGTAAATGCACAGTAATACC

15 T N V E Q D A E T N R Q C Q F T C H Y G

20 600 AACTACTTATGAGATTATGACTGTACCCAAATGACCCCTACACTTTCTCTTGTGGTGA + 660
 TTGATGAATACTCTAAACTGACATGGTTACTGGGAATGTGAAAAGAGAGAACACCACT

25 T T Y E I M T V P N D P Y T F L S C G E

30 661 AGATGGAACTGTTAGGTGGTTGATAACACGCATCAAAACTAGCTGCACAAAAGAAGATTG + 720
 TCTACCTTGACAATCCACCAAACATGTGCGTAGTTTGATCGACGTGTTCTTAAC

35 D G T V R W F D T R I K T S C T K E D C

40 721 TAAAGATGATATTTAATTAACTGTCGACGTGCTGCCACGTCTGTTGCTATTGCCACC + 780
 ATTTCTACTATAAAATTAAATTGACAGCTGACGACGGTGCAGACAACGATAAACGGGTGG

45 781 K O D I L I N C R R A A T S V A I C P P

50 AATACCATATTACCTTGTGTTGGTTGTTCTGACAGCTCAGTACGAATATATGATCGGCG + 840
 TTATGGTATAATGGAACGACAACCAACAAGACTGTCGAGTCATGCTTATATACTAGCCGC

55 I P Y Y L A V G C S D S S V R I Y D R R

60 841 AATGCTGGGACAAGAGCTACAGGGATTATGCAGGTCGAGGGACTACTGGAATGGTTGC + 900
 TTACGACCCGTGTTCTGATGTCCTTAATACGCCAGCTCCCTGATGACCTTACCAACG

65 M L G T R A T G N Y A G R G T T G M V A

70 901 CCGTTTATTCTTCCCCTTAATAATAAGTCCCTGCAGAGTGCACATCTCTGTGTTACAG + 960
 GGCAAAATAAGGAAGGGTAGAATTATTTCAGGACGTCTACTGTAGAGACACAATGTC

75 R F I P S H L N N K S C R V T S L C Y S

80 961 TGAAGATGGTCAAGAGATTCTCGTTAGTTACTCTTCAGATTACATATATCTTTTGACCC + 1020
 ACTTCTACCACTGTTCTAAGAGCAATCAATGAGAAGTCTAATGTATAGAAAAACTGGG

85 E D G Q E I L V S Y S S D Y I Y L F D P

1021 GAAAGATGATAACAGCACGAGAACCTAAAACCTCTCGCGAAGAGAGAACAGAGAGTT
CTTCTACTATGTCGTGCTCTGAATTGGAGGAAGACGCCTCTCTCTCTCTCAAA + 1080

5 K D D T A R E L K T P S A E E R R E E L

1081 GCGACAACCACCAAGTTAACGCGTTGAGACTTCGTGGTGATTGGTCAGATACTGGACCCAG
CGCTGTTGGTGGTCAATTGCAAACCTCTGAAGCACCCTAACCAAGTCTATGACCTGGTC + 1140

R O P P V K R L R L R G D W S O T G P R

1141 AGCAAGGCCGGAGAGTGAACGAGAACGAGATGGAGAGCAGAGTCCAATGTGTCAATTGAT
TCGTTCCGGCCTCTCACTTGCTCTGCTCACCTCTCGTCTCAGGGTTACACAGTAACTA + 1200

A R P E S E R E R D G E Q S P N V S L M

1201 GCAGAGAATGTCGATATGTTATCAAGATGGTTGAAGAAGCAAGTGAGGTTGCACAAAG
CGTCTCTTACAGACTATACAATAGTTCTACCAAACCTCTCGTCACTCCAACGTGTTTC + 1260

Q R M S D M L S R W F E E A S E V A O S

1261 CAATAGAGGACGAGGAAGATCTGACCCAGAGGTGGAACAAAGTCAATCAGATATTCAC + 1320

GTTATCTCCTGCTCCTCTAGAGCTGGGTCTCCACCTTGTTCAAGTTAGTCTATAAAGTTG

N R G R G R S R P R G G T S Q S D I S T

1321 TCTTCCTACGGTCCCCTCAAGTCCTGATTGGAAAGTGAGTGAAACTGCAATGGAAAGTAGA
AGAAGGATGCCAGGGTAGTTCAAGACTAAACCTTCACACTTGTACGTTACCTTCATCT + 1380

L P T V P S S P D L E V S E T A M E V D

1381 TACTCCAGCTGAACAATTCTTCAGCCTCTACATCCTCTACAATGTCAGCTCAGGCTCA
ATGAGGTCGACTTGTAAAGAAGTCGGAAAGATGTAGGAGATGTACAGTCGAGTCCGAGT + 1440

T P A E Q F L Q P S T S S T M S A Q A H

1441 TTGACATCATCTCCACAGAAAGCCCTCATTCTACTCCTTGTATCTTCAGAATAG
AAGCTGTAGTAGAGGGTGTCTTCGGGAGTAAGATGAGGAAACGATAGAAGAGGTCTATC + 1500

S T S S P T E S P H S T P L L S S P D S

1501 TGAAACAAGGCAGTCTGTTGAGGCATCTGGACACACACATCATCAGTCTGATAACAA
ACTTGTCTCCGTCAAGACAACCTCCGTAGACCTGTGGTGTGTAGTAGTCAGACTATTGTT + 1560

E Q R Q S V E A S G H H T H H Q S D N N

1561 TAATGAAAAGCTGAGCCCCAAACCAGGGACAGGTGAACCAGTTAACGTTGCAC
5 ATTACTTTCGACTCGGGGTTGGTCCCTGTCCACTTGGTCAAAATTCAAACGTGATGTC
N E K L S P K P G T G E P V L S L H Y S

1621 CACAGAAGGAACAACATAAGCACAATAAAACTGAACCTTACAGATGAATGGAGCAGTAT
10 GTGTCTTCCTTGTGATGTTCGTGTATTTGACTTGAAATGTCTACTTACCTCGTCATA
T E G T T T S T I K L N F T D E V S S I

1681 AGCATCAAGTTCTAGAGGAATTGGGAGCCATTGCAAATCTGAGGGTCAGGAGGAATCTT
15 TCGTAGTTCAAGATCTCCTTAACCCCTCGTAACGTTAGACTCCCAGTCCTCCTTAGAAA
A S S S R G I G S H C K S E G Q E E S F

1741 CGTCCCACAGAGCTCAGTGCAACCACCAAGAAGGAGACAGTGAAACAAAAGCTCCTGAAGA
20 GCAGGGTGTCTCGAGTCACGTTGGTCTTCCTCTGTCACTTGTGTTTCGAGGACTCT
V P Q S S V Q P P E G D S E T K A P E E

1801 ATCATCAGAGGATGTGACAAAATATCAGGAAGGAGTATCTGCAGAAAACCCAGTTGAGAA
25 TAGTAGTCTCCTACACTGTTTATAGTCCTCCTCATAGACGTCTTGGTCAACTCTT
S S E D V T K Y Q E G V S A E N P V E N

1861 CCATATCAATATAACACAATCAGATAAGTTCACAGCCAAGCCATTGGATTCAAACTCAGG
30 GGTATAGTTATATTGTGTTAGTCTATTCAAGTGTGGTTCGGTAACCTAACGGTTGAGTCC
H I N I T Q S D K F T A K P L D S N S G

1921 AGAAAGAAATGACCTCAATCTTGATCGCTTGTGGGTTCCAGAAGAATCTGCTTCATC
35 TCTTTCTTACTGGAGTTAGAACTAGCGAGAACACCCCAAGGTCTTCTAGACGAAGTAG
E R N D L N L D R S C G V P E E S A S S

1981 TGAAAAAGCCAAGGAACCAGAAACTTCAGATCAGACTAGCACTGAGAGTGCTACCAATGA
45 ACTTTTCGGTTCCCTGGTCTTGAAGTCTAGTCTGATCGTGAECTCAGGATGGTTACT
E K A K E P E T S D Q T S T E S A T N E

2041 AAATAACACCAATCCTGAGCCTCAGTTCAAACAGAAGCCACTGGGCCTTCAGCTCATGA
50 TTTATTGTGGTTAGGACTCGGAGTCAGGTTGTCTCGGTGACCCGGAAGTCGAGTACT
N N T N P E P O F Q T E A T G P S A H E

2101 AGAACATCCACCAGGGACTCTGCTCTTCAGGACACAGATGACAGTGATGATGACCCAGT 2160
 TCTTGTAGGTGGTCCCTGAGACGAGAAGCTCTGTCTACTGTCACTACTGGGTCA
 5 E T S T R D S A L O D T O D S O D D O P V

21-61 CCTGATCCCAGGTGCAAGGTATCGAGCAGGACCTGGTATAGACGCTCTGCTGTTGCCG 2220
 10 GGACTAGGGTCCACGTTCCATAGCTCGCTGGACCACTATCTGCAGACGACAACGGGC
 L I P G A R Y R A G P G D R R S A V A R

15 TATTCAAGGAGTTCTTCAGACGGAGAAAAGAAAGGAAAGAAATGGAAGAATTGGATACTTT 2280
 ATAAGTCCTCAAGAAGTCTGCCCTTTCTTACCTTCTTAACCTATGAAA
 20 I Q E F F R R R K E R K E M E E L D T L

2281 GAACATTAGAAGGCCGCTAGTAAAAATGGTTATAAAGGCCATCGCAACTCCAGGACAAT 2340
 CTTGTAATCTTCGGCGATCATTTTACCAAATATTCCGGTAGCGTTGAGGTCTGTTA
 25 N I R R P L V K M V Y K G H R N S R T M

2341 GATAAAAGAACCAATTCTGGGGTGCTAACCTTGTAATGACTGGTCTGAGTGTGGCCA 2400
 CTATTTCTCGGTTAAAGACCCCACGATTGAAACATTACTGACCAAGACTCACACCGGT
 30 I K E A N F W G A N F V M T G S E C G H

2401 CATTTCATCTGGGATCGGCACACTGCTGAGCATTGATGCTCTGGAAGCTGATAATCA 2460
 GTAAAAGTAGACCCCTAGCCGTGTGACGACTCGTAAACTACGAAGACCTTCGACTATTAGT
 35 I F I W D R H T A E H L M L L E A D N H

2461 TGTGGTAAACTGCCTGCAGCCACATCCGTTGACCAATTAGCCTCATCTGGCATAGA 2520
 ACACCATTTGACGGACGTGGTAGGCAAACGGTTAAATCGGAGTAGACCGTATCT
 40 V V N C L Q P H P F D P I L A S S G I D

2521 TTATGACATAAAGATCTGGTCACCATTAGAAGAGTCAGGATTAAACCGAAAATTG 2580
 AATACTGTATTTCTAGACCAAGTGGTAATCTCTCAGTTCTAAACGGTTTTGAACG
 45 Y D I K I V S P L E E S R I F N R K L A

2581 TGATGAAGTTATAACTCGAACGAACTCATGCTGGAAGAAACTAGAAACACCATTACAGT 2640
 ACTACTCAATATTGAGCTTGCTTGAGTACGACCTTCTTGATCTTGTTGTAATGTCA
 50 D E V I T R N E L M L E E T R N T I T V

2641 TCCAGCCTTTCATGTTGAGGATGTTGGCTTCACTTAATCATATCCGAGCTGACCGGTT 2700
 5 AGGTGCGAGAAAGTACAACCTACAAACCGAAGTGAATTAGTATAGGCTGACTGGCCAA
P A S F M L R M L A S L N H I R A D R L

2701 GGAGGGTGACAGATCAGAAGGCTCTGGTCAGAGAAATGAAAATGAGGATGAGGAATAATA 2760
 10 CCTCCCACGTGCTAGTCTTCCGAGACCAGTTCTCTTACTTTACTCCTACTCCTTATTAT
E G D R S E G S G Q E N E N E D E E .

15 2761 AACTCTTTGGCAAGCACTAAATGTTCTGAAATTGTATAAGACATTATTATTTTT 2820
 TTGAGAAAAACCGTTCGTGAATTACAAGACTTAAACATATTCTGAAATAATAAAAAAA

20 2821 TTTCTTACAGAGATTAGTGCATTTAACGGTTATGGTTGGAGTTTCCCTTTT 2880
 AAAGAAATGTCTCAAATCACGTTAAATTCCAATACCAAAACCTCAAAAGGGAAAAA

25 2881 TTGGGATAACCTAACATTGGTTGGAATGATTGTGTGCATGAATTGGGAGATTGTATAA 2940
 AACCTATTGGATTGTAACCAACCTTACTAACACACGTACTTAAACCTCTAACATATT

30 2941 AACAAAATAGCAGAATGTTTAAAACCTTTGCCGTGTATGAGGAGTGTAGAAAATG 3000
 TTGTTTGATCGTCTTACAAAAATTGAAACGGCACATACTCCTCACGATCTTAC

35 3001 CAAAGTGCAATATTCCTAACCTTCAAATGTGGGAGCTGGATCAATGTTGAAGAATA 3060
 GTTTCACGTTATAAAAGGGATTGAAAGTTACACCCCTGAACCTAGTTACAACCTTAT

40 3061 ATTTTCATCATAGTAAAAATGTTGGTTCAAATAAATTCTACACTTGCCTTGCATTT 3120
 TAAAAGTAGTATCCTTACACCAAGTTATTAAAGATGTGAACGGTAAACGTACAA

45 3121 TGTTGCTTCTAATTAAAGAAACTGGTTTTAAGATAACCTGAAAAAAAAAAAAAAA 3180
 ACAACGAAAGATTAATTCTTACCAACAAATTCTATGGGACTTTTTTTTTT

50 3181 AAAAAAAAAA 3189
TTTTTTTT

55 [0087] This 3164-base nucleotide sequence contains an open reading frame of 2580 nucleotides encoding Repro-EN-1.0 from nucleotide 176 to nucleotide 2755. The deduced amino acid sequence of Repro-EN-1.0 has 860 amino acids. Repro-EN-1.0 has a calculated molecular mass of 96.4 kD and a pI of 5.08.

[0088] The Repro-EN-1.0 gene encodes a 3.4 kb mRNA. This mRNA is expressed primarily in skeletal muscle, heart

and testis, and to a lesser extent in other tissues. However, it is not detected in lung or peripheral blood mononuclear cells (PBMC). Expression of Repro-EN-1.0 is up-regulated in breast and uterine carcinomas relative to their normal counterparts. It is highly expressed in both normal fallopian tube and fallopian tube carcinoma. It is expressed in low levels in normal ovary and ovarian carcinoma. Expression of the mRNA is lower in endometrial carcinoma cell lines than in prostate adenocarcinoma cell lines.

5 [0089] Analysis of the deduced amino acid sequence of Repro-EN-1.0 shows no significant sequence identity with any other protein.

[0090] There is an alternately spliced variant that was isolated from a human heart cDNA library. This variant is called IB1 and is useful in the same ways as the nucleotide sequence (SEQ ID NO:1) and deduced amino acid sequence (SEQ ID NO:2) of Repro-EN-1.0. The IB1 sequence was isolated from a heart cDNA library by screening with a nucleic acid probe obtained from the Repro-EN-1.0 sequence. The IB1 sequence is different from the Repro-EN-1.0 sequence in that it contains an additional 231 bp exon inserted into the cDNA sequence at position 1555. Therefore the IB1 sequence has similar properties, but is slightly larger.

[0091] The nucleotide sequence (SEQ ID NO:3) and the deduced amino acid sequence (SEQ ID NO:4) of IB1 follow:

15

1 CGGCCGGGCTTCAGGGGCCAGGCGCCGCTGCTGCCACCGCCATCTAACGCTGCGCCCTG
GCCGGCCCCAAGTCCCCGGGTCCCGGGCGACGACGGTGGCGGTAGATTGCGACGCCGGGAC 60

20

61 GAGGCCCGGCAGCGATGGTGCCGGTGCAGCTCGGGTGTGAAACGGGTGTCCTCTCCC
CTCCGGGCCGCGCCTACACGGCACGCCAGGCCACAACTTGCCACAGGGGAGGG 120

25

121 CCTCCTCCCCCTCCCCCACGGGTGGTCTCCCTCCACCCGGCTCAGGCAGAGCCATGTC
GGAGGAGGGAGGGGGTGCGCCACCAAGAGGGAGGGTGGGCCAGTCCGTCTCGGTACAG 180

30

M S

35

40

45

50

55

181 TCGGGGTGGCTCCTACCCACACCTGTTGGACGTGAGGAAAAGGTTCTCGGGCTGGA
 + 240
 AGCCCCACCGAGGATGGGTGTGGACAACACCTTGCACTCTTTCCAAGGAGCCCACCT
 5 R G G S Y P H L L W D V R K R F L G L E

 241 GGACCCGTCGGCTCGGGAGTCGCTACCTGGAAAGAAGAGAATTATCAAAGATTAAA
 + 300
 CCTGGCAGGGCCGACGCCCTAGCGATGGACCCCTTCTCTTAATAGGTTCTAATTT
 10 D P S R L R S R Y L G R R E F I Q R L K

 301 ACTTGAAGCAACCCTTAATGTGCATGATGGTTGTGTTAACATAACTGTTGGAATGACAC
 + 360
 TGAACCTCGTGGATTACACGTACTACCAACACAATTATGTTAGACAACCTTACTGTG
 15 L E A T L N V H D G C V N T I C W N D T

 361 TGGAGAAATATTTTATCTGGCTCAGATGACACCAAAATAGTAAATAGTAATCCTTACAG
 + 420
 ACCTCTTATATAAAATAGACCGAGTCTACTGTGGTTAACATTAAATCATTAGGAATGTC
 20 G E Y I L S G S D D T K L V I S N P Y S
 25 CAGAAAGGTTTGACAACAATTGTTCAAGTCCCCTGGCTCGTTGTATAAATCACGTTCAA
 + 480
 GTCTTCAAAACTGTTAGCAAGTCCCCTGGCTCGTTGTATAAATCACGTTCAA
 30 R K V L T T I R S G H R A N I F S A K F

 481 CTTACCTTGTACAAATGATAAACAGATTGTATCCTGCTCTGGAGATGGAGTAATATTTA
 + 540
 GAATGGAACATGTTACTATTTGCTAACATAGGACGAGACCTCACCTATTATAAAAT
 35 L P C T N D K Q I V S C S G D G V I F Y

 541 TACCAACGTTGAGCAAGATGCAGAAACCAACAGACAATGCCAATTACGTGTCATTATGG
 + 600
 ATGGTTGCAACTCGTTACGTCTTGGTTGTCTACGGTTAACATGACAGTAATACC
 40 T N V E Q D A E T N R Q C Q F T C H Y G

 601 AACTACTTATGAGATTATGACTGTACCCAAATGACCCCTTACACTTTCTCTTGTGGTGA
 + 660
 TTGATGAATACTCTAACATGGTTACTGGGAATGTGAAAAGAGAGAACACCACT
 45 T T Y E I M T V P N D P Y T F L S C G E

 661 AGATGGAACGTAGGTGGTTGATAACCGCATCAAAACTAGCTGCACAAAAGAAGATTG
 + 720
 TCTACCTTGACAATCCACCAAACATATGTCGTAGTTTGATCGACGTGTTCTTCAAC
 50 D G T V R W F D T R I K T S C T K E D C

721 TAAAGATGATATTTAATTAACGTGCGACGTGCTGCCACGTCTGTTGCTATTGCCACC
 780 ATTTCTACTATAAAATTAGACAGCTGCACGACGGTGCAGACAACGATAAACGGGTGG
 5 K D D I L I N C R R A A T S V A I C P P

 781 AATACCATATTACCTTGCTGTTGGTTCTGACAGCTAGTACGAATATATGATCGGCG
 840 TTATGGTATAATGGAACGACAACCAACAAGACTGTCGAGTCATGCTTATATACTAGCCGC
 10 I P Y Y L A V G C S D S S V R I Y D R R

 841 AATGCTGGGCACAAGAGCTACAGGGATTATGCAGGTGAGGGACTACTGGAATGGTTGC
 900 TTACGACCCGTGTTCTCGATGTCCCTTAATACGTCCAGCTCCCTGATGACCTTACCAACG
 15 M L G T R A T G N Y A G R G T T G M V A

 901 CCGTTTTATTCTCTCCATCTTAATAAAAGTCCTGCAGAGTGACATCTCTGTTACAG
 960 GGCAAAATAAGGAAGGGTAGAATTATTATTAGGACGTCTCACTGTAGAGACACAATGTC
 20 R F I P S H L N N K S C R V T S L C Y S
 25
 961 TGAAGATGGTCAAGAGATTCTCGTTAGTTACTCTTCAGATTACATATATCTTTGACCC
 1020 ACTTCTACCACTGCTTAAGAGCAATCAATGAGAAGTCAATGTATAGAAAAACTGGG
 30 E D G Q E I L V S Y S S D Y I Y L F D P

 1021 GAAAGATGATACAGCACGAGAACTAAAACCTCTGCAGAGAGAAGAGAGTT
 1080 CTTTCTACTATGTCGTGCTCTGAATTGGAGAGACGCCCTCTCTCTCTCAA
 35 K D D T A R E L K T P S A E E R R E E L

 1081 GCGACACCACCAAGTTAACGCGTTGAGACTTCGTGGTATTGGTCAGATACTGGACCCAG
 1140 CGCTGTTGGTGGTCAATTGCAAAACTCTGAAGCACCCTAACCAAGTCTATGACCTGGGTC
 40 R Q P P V K R L R L R G D W S D T G P R

 1141 AGCAAGGCCGGAGAGTGAACGAGAACGAGAGATGGAGAGCAGAGTCCCAATGTGTCAATTGAT
 1200 TCGTTCCGGCTCTCACTTGCTCTGCTCACCTCTCGTCTCAGGGTACACAGTAACTA
 45 A R P E S E R E R O G E Q S P N V S L M

 1201 GCAGAGAAATGTCGATATGTTATCAAGATGGTTGAAGAAGCAAGTGAGGTTGCACAAAG
 1260 CGTCTCTTACAGACTATACAATAGTTCTACCAAACTCTTCGTTCACTCCAACGTGTTTC
 50 Q R M S D M L S R W F E E A S E V A O S

1261 CAATAGAGGACGAGGAAGATCTGACCCAGAGGTGAAACAAGTCATCAGATATTC AAC
GTTATCTCCTGCTCCCTCTAGAGCTGGGCTCCACCTTGTTCAGTTAGTCTATAAAGTTG
5 N R G R G R S R P R G G T S Q S D I S T

1321 TCTTCCTACGGTCCCCTCAAGTCCTGATTGAAAGTGAGTGAAACTGCAATGAAAGTAGA
AGAAGGATGCCAGGGTAGTTCAAGGACTAACCTTCACTCACTTGACGTTACCTTCATCT
10 L P T V P S S P D L E V S E T A M E V D

1381 TACTCCAGCTGAACAATTCTTCAGCCTTACATCCTCTACAATGTCAGCTCAGGCTCA
ATGAGGTCGACTTGTAAAGAAGTCGGAAAGATGTAGGAGATGTTACAGTCGAGTCCGAGT
15 T P A E Q F L Q P S T S S T M S A Q A H

1441 TTGACATCATCTCCCACAGAAAGCCCTCATTCTACTCCTTGCTATCTTCAGATAG
AAGCTGTAGTAGAGGGTGTCTTCGGGAGTAAGATGAGGAAACGATAGAAGAGGTCTATC
20 S T S S P T E S P H S T P L L S S P D S

1501 TGAAACAAAGGCAGTCTGTTGAGGCATCTGGACACCACACATCATCAGTCTGAATT
ACTTGTTCCGTCAAGACAACCTCCGTAGACCTGTGGTGTGTAGTAGTCAGACTAAAAAA
25 E Q R Q S V E A S G H H T H H Q S E F L

1561 AAGGGGGCCTGAGATAGCTTCTCGTAAGCGCTGCAACAACTGAGGCTTAAGAAGGC
TTCCCCCGGACTCTATCGAAACGAAGCATTGCGGACGTTGACTCCGAATTCTTCCG
30 R G P E I A L L R K R L Q Q L R L K K A

231 bp insert

1621 TGAGCAGCAGAGGGCAGCAAGAGCTAGCTCACATACCCAGCAACAGCCTTCACTTC
ACTCGTCGTCTCGTCGTTCTCGATCGACGTGTATGGTCGTTGAGACT
35 E Q Q R Q Q E L A A H T Q Q Q P S T S D

231 bp insert

1681 TCAGTCTTCTCATGAGGGCTTCAACAGGACCCATGCTTCAGATTCTCCTTCTGT
AGTCAGAAGAGTACTCCGAGAAGTGTCTGGGAGTACGAAGTCTAAGAGGAAGACA
40 Q S S H E G S S Q D P H A S D S P S S V

231 bp insert

45

50

55

1741 GGTTAACAAACAGCTCGGATCCATGTC ACTTGACGAGCAACAGGATAACAATAATGAAAAA
 CCAATTGTTTGTGAGCCTAGGTACAGTGAACTGCTCGTTGCCATTGTTATTACTTTT + 1800

5 V N K Q L G S M S L D E Q Q D N N N E K

 231 bp insert →

1801 GCTGAGCCCCAAACCAGGGACAGGTGAACCAAGTTTAAGTTGCAC TACAGCACAGAAGG
 CGACTCGGGTTGGTCCCTGTCCACTTGGTCAAACGTGATGTCGTGCTTCC + 1860

10 L S P K P G T G E P V L S L H Y S T E G

1861 AACAACTACAAGCACAAATAAAACTGAACCTTACAGATGAATGGAGCAGTATAGCATCAAG
 TTGTTGATGTTCGTGTATTGACTTGAAATGTCTACTTACCTCGTCATATCGTAGTTC + 1920

15 T T T S T I K L N F T O E W S S I A S S

1921 TTCTAGAGGAATTGGGAGCCATTGCAAATCTGAGGGTCAGGAGGAATCTTCGTCCCACA
 AAGATCTCCTTAACCCCTCGGTAACGTTAGACTCCCAGTCCTCCCTAGAAAGCAGGGTGT + 1980

20 S R G I G S H C K S E G Q E E S F V P Q

1981 GAGCTCAGTGCAACCACCAAGAAGGAGACAGTGAAACAAAAGCTCCTGAAGAATCATCAGA
 CTCGAGTCACGTTGGTGGTCTTCCTCTGTCACTTGTGTTCGAGGACTTCTTAGTAGTCT + 2040

25 S S V Q P P E G D S E T K A P E E S S E

2041 GGATGTGACAAAATATCAGGAAGGAGTATCTGCAGAAAACCCAGTTGAGAACCATATCAA
 CCTACACTGTTTATAGTCCTCCTCATAGACGTCTTGGTCAACTCTGGTATAGTT + 2100

30 D V T K Y Q E G V S A E N P V E N H I N

2101 TATAACACAATCAGATAAGTTCACAGCCAAGCCATTGGATTCCAACTCAGGAGAAAGAAA
 ATATTGTGTTAGTCTATTCAAGTGTGGTCCGGTAACCTAAGGTTGAGTCCTCTTCTT + 2160

35 I T Q S D K F T A K P L D S N S G E R N

2161 TGACCTCAATTTGATCGCTTGTGGGTTCCAGAAGAATCTGCTTCATCTGAAAAAGC
 ACTGGAGTTAGAACTAGCGAGAACACCCCAAGGTCTTAGACGAAGTAGACTTTTCG + 2220

40 D L N L D R S C G V P E E S A S S E K A

2221 CAAGGAACCAGAAAATTCAAGACTAGCACTGAGAGTGCTACCAATGAAAATAACAC
 GTTCTTGGTCTTGAAGTCTAGTCTGACTCTCACGATGGTTACTTTATTGTG + 2280

45 K E P E T S D Q T S T E S A T N E N N T

2281 CAAT CCTGAG CCTCAG TTCCAAAC AGAAGCCACTGGGCCTTCAGCTCATGAAGAACATC
5 GTTGGAGACTCGGAGTCAGGTTGTCTCGGTGACCCGGAAGTCGAGTACTTCTTTGTAG
N P E P Q F O T E A T G P S A H E E T S

2341 CACCAGGGACTCTGCTCTTCAGGACACAGATGACAGTGATGATGACCCAGTCCTGATCCC
10 GTGGTCCCTGAGACGAGAAGTCTGTCTACTGTCACTACTGGGTAGGACTAGGG
T R D S A L Q D T D D S D D D P V L I P

2401 AGGTGCAAGGTATCGAGCAGGACCTGGTGTAGACGCTCTGCTGTTGCCGTATTCAAGGA
15 TCCACGTTCCATAGCTCGTCCTGGACCACTATCTGCAGACGACAACGGGCATAAGTCCT
G A R Y R A G P G D R R S A V A R I Q E

2461 GTTCTTCAGACGGAGAAAAGAAAGGAAAGAAATGGAAGAATTGGATACTTTGAACATTAG
20 CAAGAAGTCTGCCCTTTCTTACCTTAACTGAAACTTGTAAATC
F F R R R K E R K E M E E L D T L N I R

2521 AAGGCCGCTAGTAAAATGGTTATAAAGGCCATCGCAACTCCAGGACAATGATAAAAGA
25 CTCGGCGATCATTTACCAAATATTCGGTAGCGTTGAGGTCTGTTACTATTTCT
R P L V K M V Y K G H R N S R T M I K E

2581 AGCCAATTCTGGGTGCTAACCTTGTAATGACTGGTCTGAGTGTGCCACATTTCAT
30 TCGGTTAAAGACCCACGATTGAAACATTACTGACCAAGACTCACACCGGTAAAAGTA
A N F W G A N F V M T G S E C G H I F I

2641 CTGGGATCGGCACACTGCTGAGCATTGATGCTTCTGGAAAGCTGATAATCATGTGGTAAA
35 GACCCTAGCCGTGACGACTCGTAAACTACGAAGACCTTCGACTATTAGTACACCATT
W D R H T A E H L M L L E A D N H V V N

2701 CTGCCTGCAGCCACATCCGTTGACCAATTAGCTCATCTGGCATAGATTATGACAT
40 GACGGACGTGGTAGGCAAACGGTTAAAATCGGAGTAGACCGTATCTAAACTGTA
C L Q P H P F D P I L A S S G I D Y D I

2761 AAAGATCTGGTCACCATAGAAGAGTCAGGATTTAACCGAAAACCTGCTGATGAAGT
45 TTTCTAGACCAAGTGGTAAATCTCTCAGTTCTAAACGGCTTTGAACGACTACTTCA
K I W S P L E E S R I F N R K L A D E V

2821 TATAACTCGAAACGAACATGCTGGAAGAAACTAGAAAACCATTACAGTTCCAGCCTC 2880
 ATATTGAGCTTGCTTGAGTACGACCTCTTGATCTTGTGGTAATGTCAAGGTCGGAG
 5 I T R N E L M L E E T R N T I T V P A S

2881 TTTCATGTTGAGGATGTTGGCTTCACTTAATCATATCCGAGCTGACCGGTTGGAGGGTGA 2940
 AAAGTACAACCTCCTACAAACCGAAGTGAATTAGTATAGGCTCGACTGCCAACCTCCACT
 10 F M L R M L A S L N H I R A D R L E G D

2941 CAGATCAGAAGGGCTCTGGTCAAGAGAAATGAAAATGAGGATGAGGAATAATAAACTCTTT 3000
 GTCTAGTCTTCCGAGACCAGTTCTCTTACTTTACTCCTACTCCTTATTATTTGAGAAAA
 15 R S E G S G Q E N E N E D E E .

3001 TGGCAAGCACTAAATGTTCTGAAATTGTATAAGACATTATTATTTTTTTCTTAC 3060
 ACCGTTCGTGAATTACAAGACTTAAACATATTCTGAAATAATAAAAAAAAAGAAATG
 20

3061 AGAGATTAGTGCAATTAAAGGTTATGGTTGGAGTTTCCCTTTGGATAA 3120
 TCTCTAAATCACGTTAAATTCCAATACCAAAACCTCAAAAGGGAAAAACCTATT
 25

3121 CCTAACATTGGTTGGAATGATTGTGTGCATGAATTGGAGATTGTATAAAACAAAAC 3180
 GGATTGTAACCAAACCTTACTAACACACGTACTTAAACCCCTCTAACATATTTGTTTGA
 30

3181 AGCAGAAATGTTTAAACCTTTGCCGTGTATGAGGAGTGCTAGAAAATGCAAAGTCA 3240
 TCGCTTACAAAAATTGAAAAACGGCACATACTCCTCACGATTTACGTTCACGT
 35

3241 ATATTTCCCTAACCTCAAATGTGGAGCTGGATCAATGTTGAAGAATAATTTCATC 3300
 TATAAAAGGGATTGGAAGTTACACCCCTCGAACCTAGTTACAACCTTCTTAAAGTAG
 40

3301 ATAGTAAAAATGTTGGTTCAAATAAATTCTACACTTGCCTTGATGTTGTTGCTT 3360
 TATCACTTTACAACCAAGTTATTAAAGATGTGAACGGTAAACGTACAAACAACGAAA
 45

3361 CTAATTAAGAAACTGGTTTTAAGATACCCCTGAAAAAAAAAAAAAAA 3420
 GATTAATTCTTGACCAACAAAATTCTATGGACTTTTTTTTTTTTTTTTTTT
 50

IB1 Alternately Spliced Variant from Heart Library:

- 55
 IB1 cDNA is 3,395 base pairs
 IB1 open reading frame is 2,811 nucleotides
 IB1 open reading frame maps to nucleotides 176 to 2986

Deduced amino acid sequence of IB1 has 937 amino acids
 Calculated molecular mass of IB1 protein is 104,969 Daltons (105kD)
 Calculated pI of IB1 protein is 5.17

- 5 [0092] Analysis of the amino acid sequence identified several amino acid motifs that will be apparent to those skilled in the art including a myb 1 DNA binding domain, a WD 40 site, an RGD cell-attachment sequence, an N-myristylation site and several phosphorylation and glycosylation sites. Analysis also shows that the protein is largely hydrophilic. This implies that most of the amino acid sequence is exposed to the immune system and can be recognized as epitopes.
- 10 [0093] Analysis of expressed sequence tags (ESTs) from a public database (Genbank) identified many overlapping ESTs that, together, covered most of the Repro-EN-1.0 cDNA sequence.

III. REPRO-EN-1.0 and IB1 NUCLEIC ACIDS

15 [0094] This invention provides recombinant polynucleotides comprising a nucleotide sequence encoding Repro-EN-1.0 and IB1 proteins, Repro-EN-1.0 and IB1 analogs or fragments of these polypeptides, as described herein. Repro-EN-1.0 or IB1 analogs include 1) immunogenic fragments of Repro-EN-1.0 or IB1; 2) homologs of Repro-EN-1.0 or IB1 from other mammals (especially primates); 3) fragments of Repro-EN-1.0 or IB1 comprising at least 5 consecutive amino acids from the sequence of Repro-EN-1.0 or IB1; 4) non-naturally occurring polypeptides whose sequences are substantially identical to Repro-EN-1.0; and 5) fusion proteins comprising a Repro-EN-1.0 or IB1 or a Repro-EN-1.0 or IB1 analog fused to a second polypeptide moiety. The polynucleotides are useful for expressing the mRNA or polypeptides they encode and in the preparation of probes or primers, among other things.

20 [0095] In one embodiment, the recombinant polynucleotide molecule comprises a nucleotide sequence encoding a sequence of at least 5 amino acids selected from the amino acid sequence of Repro-EN-1.0 (SEQ ID NO:2) or IB (SEQ ID NO:4). The nucleotide sequence can encode a sequence of at least 25 amino acids, at least 100 amino acids or at least 200 amino acids from SEQ ID NO:2 or SEQ ID NO:4. In one embodiment, the nucleotide sequence encodes an immunogenic analog. One such immunogenic analog is a polypeptide comprising an epitope that binds specifically to an antibody from serum from a subject diagnosed with endometriosis. In another embodiment, the nucleotide sequence encodes full-length native Repro-EN-1.0 or IB1 polypeptide.

25 [0096] The nucleotide sequence can be identical to a sequence from Repro-EN-1.0 cDNA or its complement or IB1 cDNA or its complement, or can include degenerate codons. In one embodiment of a nucleotide sequence encoding full-length Repro-EN-1.0 or IB1, the sequence is identical to the coding sequence of Repro-EN-1.0 of SEQ ID NO:1 or IB1 of (SEQ ID NO:3). In another embodiment, the nucleotide sequence encodes a Repro-EN-1.0 or IB1 analog whose amino acid sequence is substantially identical to the amino acid sequence of Repro-EN-1.0 polypeptide (SEQ ID NO:2) or IB1 polypeptide (SEQ ID NO:4).

30 [0097] In another embodiment, the polynucleotide encodes a fusion protein between Repro-EN-1.0 or IB1 polypeptide or Repro-EN-1.0 or IB1 analog amino acid sequences and a second amino acid sequence. The second amino acid sequence can be, for example, a detectable label such as a fluorescent protein, enzyme marker of protein from a two-hybrid system.

35 [0098] The polynucleotides of the present invention are cloned or amplified by *in vitro* methods, such as the polymerase chain reaction (PCR), the ligase chain reaction (LCR), the transcription-based amplification system (TAS), the self-sustained sequence replication system (3SR) and the Q β replicase amplification system (QB). For example, a polynucleotide encoding the protein can be isolated by polymerase chain reaction of cDNA from a human endometrial carcinoma cell line using primers based on the DNA sequence of Repro-EN-1.0 of SEQ ID NO:1. One pair of primers useful for amplifying Repro-EN-1.0 DNA, including allelic variants, is:

40 45 Upstream sense: 5'-caggacacagatgacagtgt-3' (SEQ ID NO:5)
 Downstream antisense: 5'-agaggccctctgtatctgtcac-3' (SEQ ID NO:6).

50 [0099] A wide variety of cloning and *in vitro* amplification methodologies are well-known to persons of skill. PCR methods are described in, for example, U.S. Pat. No. 4,683,195; Mullis et al. (1987) *Cold Spring Harbor Symp. Quant. Biol.* 51:263; and Erlich, ed., *PCR Technology*, (Stockton Press, NY, 1989). One useful format is real time PCR. See, e.g., Luch et al. (1997) *J. Molec. Endocrinol.* 18:77-85 and Arold et al., (1997) *Proc. Nat'l Acad. Sci., USA* 94:2438-43. Polynucleotides also can be isolated by screening genomic or cDNA libraries with probes selected from the sequences of SEQ ID NO:1 under stringent hybridization conditions.

55 [0100] Mutant versions of the proteins can be made by site-specific mutagenesis of other polynucleotides encoding the proteins, or by random mutagenesis caused by increasing the error rate of PCR of the original polynucleotide with 0.1 mM MnCl₂ and unbalanced nucleotide concentrations.

[0101] This invention also provides expression vectors, e.g., recombinant polynucleotide molecules comprising ex-

pression control sequences operatively linked to a nucleotide sequence encoding the target polypeptide. Expression vectors can be adapted for function in prokaryotes or eukaryotes by inclusion of appropriate promoters, replication sequences, markers, etc. for transcription and translation of mRNA. The construction of expression vectors and the expression of genes in transfected cells involves the use of molecular cloning techniques also well known in the art.

5 Sambrook et al., *Molecular Cloning -- A Laboratory Manual*, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY, (1989) and *Current Protocols in Molecular Biology*, F.M. Ausubel et al., eds., (Current Protocols, a joint venture between Greene Publishing Associates, Inc. and John Wiley & Sons, Inc.) Useful promoters for such purposes include a metallothionein promoter, a constitutive adenovirus major late promoter, a dexamethasone-inducible MMTV promoter, a SV40 promoter, a MRP polIII promoter, a constitutive MPSV promoter, a tetracycline-inducible CMV promoter (such as the human immediate-early CMV promoter), and a constitutive CMV promoter.

10 [0102] Methods for transfecting genes into mammalian cells and obtaining their expression for *in vitro* use or for gene therapy, are well known to the art. See, e.g., *Methods in Enzymology*, vol. 185, Academic Press, Inc., San Diego, CA (D.V. Goeddel, ed.) (1990) or M. Krieger, *Gene Transfer and Expression -- A Laboratory Manual*, Stockton Press, New York, NY, (1990).

15 [0103] Expression vectors useful in this invention depend on their intended use. Such expression vectors must, of course, contain expression and replication signals compatible with the host cell. Expression vectors useful for expressing the protein of this invention include viral vectors such as alpha viruses, retroviruses, adenoviruses and adeno-associated viruses, plasmid vectors, cosmids, liposomes and the like. Viral and plasmid vectors are preferred for transfecting mammalian cells. The expression vector pcDNA1 (Invitrogen, San Diego, CA), in which the expression control sequence comprises the CMV promoter, provides good rates of transfection and expression. Adeno-associated viral vectors are useful in the gene therapy methods of this invention.

20 [0104] The construct can also contain a tag to simplify isolation of the protein. For example, a polyhistidine tag of, e.g., six histidine residues, can be incorporated at the amino terminal end of the protein. The polyhistidine tag allows convenient isolation of the protein in a single step by nickel-chelate chromatography.

25 [0105] In another embodiment, endogenous genes are transcribed by operatively linking them to expression control sequences supplied endogenously that recombine with genomic DNA. In one method, one provides the cell with a recombinant polynucleotide containing a targeting sequence, which permits homologous recombination into the genome upstream of the transcriptional start site of target gene; the expression control sequences; an exon of the target gene; and an unpaired splice-donor site which pairs with a splice acceptor in the target gene. Such methods are discussed in Treco et al., WO 94/12650; Treco et al., WO 95/31560 and Treco et al., WO 96/29411.

30 [0106] The invention also provides recombinant cells comprising an expression vector for expression of the nucleotide sequences encoding a polypeptide of this invention. Host cells can be selected for high levels of expression in order to purify the protein. Mammalian cells are preferred for this purpose, but prokaryotic cells, such as *E. coli*, also are useful. The cell can be, e.g., a recombinant cell in culture or a cell *in vivo*.

35

IV. POLYNUCLEOTIDE PROBES AND PRIMERS

40 [0107] This invention provides polynucleotide probes and primers that specifically hybridize to a sub-sequence of Repro-EN-1.0 cDNA or its complement or IB1 cDNA or its complement, under stringent hybridization conditions. The probes and primers of this invention are polynucleotides of at least 7 nucleotides, at least 10 nucleotides, at least 15 nucleotides, at least 20 nucleotides or at least 25 nucleotides. In one embodiment, the sequence of the polynucleotide is a contiguous sequence from SEQ ID NO: 1 or its complement. Any suitable region of the Repro-EN-1.0 or IB1 gene may be chosen as a target for polynucleotide hybridization. Nucleotide substitutions, deletions, and additions may be incorporated into the polynucleotides as long as the characteristic ability to specifically hybridize to the target sequence or its complement is retained. Nucleotide sequence variation may result from sequence polymorphisms of various alleles, minor sequencing errors, and the like.

45 [0108] The probes and primers of the invention are useful as probes in hybridization assays, such as Southern and northern blots, for identifying polynucleotides having a nucleotide sequence encoding a Repro-EN-1.0 or IB1 polypeptide, and as primers for amplification procedures. The probes and primers of the invention are also useful in detecting the presence, absence or amount of Repro-EN-1.0 or IB1 in tissue biopsies and histological sections where the detection method is carried out *in situ*, typically after amplification of Repro-EN-1.0 or IB1 sequences using a primer set.

50 [0109] The probes and primers of this invention also are useful for identifying allelic forms of Repro-EN-1.0 and animal cognate genes or IB1 and animal cognate genes. Probes and primers can be used to screen human or animal genomic DNA or cDNA libraries under, e.g., stringent conditions. DNA molecules that specifically hybridize to the probe are then further examined to determine whether they are Repro-EN-1.0 allelic variants or animal cognates or IB1 allelic variants or animal cognates.

55 [0110] The probes also are useful in oligonucleotide arrays. Such arrays are used in hybridization assays to check the identity of bases in a target polynucleotide. In essence, when a target hybridizes perfectly to a probe on the array,

the target contains the nucleotide sequence of the probe. When the target hybridizes less well, or does not hybridize at all, then the target and probe differ in sequence by one or more nucleotide. By proper selection of probes, one can check bases on a target molecule. See, e.g., Chee et al., WO 95/11995. The use the Repro-EN-1.0 or IB1 sequence in genomics is described further below.

5 [0111] In one embodiment, the polynucleotide is directly or indirectly detectable through a detectable moiety. A detectable moiety bound to either an oligonucleotide primer or a probe is subsequently used to detect hybridization of an oligonucleotide primer to the RNA component. Detection of labeled material bound to a Repro-EN-1.0 or IB1 polynucleotide in a sample provides a means of determining a diagnostic or prognostic value.

10 [0112] Although primers and probes can differ in sequence and length, the primary differentiating factor is one of function: primers serve as an initiation point for DNA synthesis of a target polynucleotide, as in RT and PCR reactions, while probes are typically used for hybridization to and detection of a target polynucleotide. Typical lengths of primers or probes can range from 7-50 nucleotides, preferably from 10-40 nucleotides, and most preferably from 15-35 nucleotides. A primer or probe can also be labeled with a detectable moiety for detection of hybridization of the primer or probe to the target polynucleotide.

15 [0113] In general, those of skill in the art recognize that the polynucleotides used in the invention include both DNA and RNA molecules and naturally occurring modifications thereof, as well as synthetic, non-naturally occurring analogs of the same, and heteropolymers, of deoxyribonucleotides, ribonucleotides, and/or analogs of either. The particular composition of a polynucleotide or polynucleotide analog will depend upon the purpose for which the material will be used and the environment in which the material will be placed. Modified or synthetic, non-naturally occurring nucleotides have been designed to serve a variety of purposes and to remain stable in a variety of environments, such as those in which nucleases are present.

20 [0114] Oligonucleotides preferably are synthesized, e.g., on an Applied BioSystems or other commercially available oligonucleotide synthesizer according to specifications provided by the manufacturer. Oligonucleotides may be prepared using any suitable method, such as the phosphotriester and phosphodiester methods, or automated embodiments thereof. In one such automated embodiment, diethylphosphoramidates are used as starting materials and may be synthesized as described by Beaucage et al., *Tetrahedron Letters* 22: 1859 (1981), and U.S. Patent No. 4,458,066.

25 [0115] Polynucleotides, e.g., probes, also can be recombinantly produced through the use of plasmids or other vectors.

30 [0116] In one aspect this invention provides a probe that specifically hybridizes to the 5' untranslated region of Repro-EN-1.0 or IB1, the coding region of Repro-EN-1.0 or IB1, or a region of Repro-EN-1.0 or IB1 encoding an epitope of the Repro-EN-1.0 or IB1 polypeptide.

35 [0117] In another aspect, this invention provides a primer pair which amplifies a nucleotide sequence encoding a polypeptide epitope of Repro-EN-1.0 or IB1 recognized by an antibody from an individual diagnosed with endometriosis. A primer pair that amplifies a particular nucleotide sequence (given in the 5' to 3' orientation) includes a 5' primer and a 3' primer. The 3' primer hybridizes to the 3' end of the nucleotide sequence or downstream from it. The 5' primer hybridizes to the 3' end of the complement of the nucleotide sequence or downstream from it. In this way, the primers can amplify a polynucleotide that comprises the nucleotide sequence. One nucleotide sequence encoding a polypeptide epitope of Repro-EN-1.0 has been identified within the about 2.2 kb from 3' end of the coding sequence of Repro-EN-1.0 (SEQ ID NO:1).

40 V. METHODS FOR DETECTING REPRO-EN-1.0 and IB1 POLYNUCLEOTIDES

[0118] The probes and primers of this invention are useful, among other things, in detecting Repro-EN-1.0 or IB1 polynucleotides in a sample. A method for detecting the presence, absence or amount of a Repro-EN-1.0 or IB1 polynucleotide in a sample involves two steps: (1) specifically hybridizing a polynucleotide probe or primer to a Repro-EN-1.0 or IB1 polynucleotide, and (2) detecting the specific hybridization.

45 [0119] For the first step of the method, the polynucleotide used for specific hybridization is chosen to hybridize to any suitable region of Repro-EN-1.0. The polynucleotide can be a DNA or RNA molecule, as well as a synthetic, non-naturally occurring analog of the same. The polynucleotides in this step are polynucleotide primers and polynucleotide probes disclosed herein. This includes probes and primers having sequences selected from the sequence of Repro-EN-1.0 (SEQ ID NO:1) or selected from the cyber-sequence of Repro-EN-1.0 (SEQ ID NO:5).

50 [0120] For the second step of the reaction, any suitable method for detecting specific hybridization of a polynucleotide to Repro-EN-1.0 or IB1 may be used. Such methods include, e.g., amplification by extension of a hybridized primer using reverse transcriptase (RT); extension of a hybridized primer using RT-PCR or other methods of amplification; and *in situ* detection of a hybridized primer. In *in situ* hybridization, a sample of tissue or cells is fixed onto a glass slide and permeabilized sufficiently for use with *in situ* hybridization techniques. Detectable moieties used in these methods include, e.g., labeled polynucleotide probes; direct incorporation of label in amplification or RT reactions, and labeled polynucleotide primers.

[0121] Often, cell extracts or tissue samples used in methods for determining the amount of a polynucleotide in a sample will contain variable amounts of cells or extraneous extracellular matrix materials. Thus, a method for determining the cell number in a sample is important for determining the relative amount per cell of a test polynucleotide such as Repro-EN-1.0 or IB1. A control for cell number and amplification efficiency is useful for determining diagnostic values for a sample of a potential cancer, and a control is particularly useful for comparing the amount of test polynucleotide such as Repro-EN-1.0 or IB1 in sample to a diagnostic value for breast cancer, uterine cancer or fallopian tube cancer. A preferred embodiment of the control RNA is endogenously expressed 28S rRNA. (See, e.g., Khan *et al.*, *Neurosci. Lett.* 147: 114-117 (1992) which used 28S rRNA as a control, by diluting reverse transcribed 28S rRNA and adding it to the amplification reaction.)

VI. INHIBITORY POLYNUCLEOTIDES FOR INHIBITING REPRO-EN-1.0 AND IB1 EXPRESSION

A. General

[0122] This invention also provides inhibitory polynucleotides directed against Repro-EN-1.0 or IB1 polynucleotides that inhibit Repro-EN-1.0 or IB1 expression and, therefore inhibit its activity in a cell. Inhibitory polynucleotides can inhibit Repro-EN-1.0 or IB1 activity in a number of ways. According to one mechanism, the polynucleotide prevents transcription of the Repro-EN-1.0 or IB1 gene (for instance, by triple helix formation). In another mechanism, the polynucleotide destabilizes the Repro-EN-1.0 or IB1 and reduces its half-life. In another mechanism, the polynucleotide inhibits assembly of the RNA component into the Repro-EN-1.0 or IB1 by binding to Repro-EN-1.0 or IB1.

[0123] An inhibitory polynucleotide is a polynucleotide that is capable of specifically hybridizing with a target polynucleotide and that interferes with the transcription, processing, translation or other activity the target polynucleotide. Inhibitory polynucleotides generally are single-stranded and have a sequence of at least 7, 8, 9, 10, or 11 nucleotides capable of specifically hybridizing to the target sequence. RNA sequences generally require a sequence of at least 10 nucleotides for specific hybridization. Inhibitory polynucleotides include, without limitation, antisense molecules, ribozymes, sense molecules and triplex-forming molecules. In one embodiment, the inhibitory polynucleotide is no more than about 50 nucleotides long.

[0124] While not wishing to be limited by theory, it is believed that inhibitory polynucleotides inhibit the function of a target, in part, by binding to the appropriate target sequence. An inhibitory polynucleotide can inhibit DNA replication or DNA transcription by, for example, interfering with the attachment of DNA or RNA polymerase to the promoter by binding to a transcriptional initiation site or a template. It can interfere with processing of mRNA, poly(A) addition to mRNA or translation of mRNA by, for example, binding to regions of the RNA transcript such as the ribosome binding site. It can promote inhibitory mechanisms of the cells, such as promoting RNA degradation via RNase action. The inhibitory polynucleotide can bind to the major groove of the duplex DNA to form a triple helical or "triplex" structure. Methods of inhibition using inhibitory polynucleotides therefore encompass a number of different approaches to altering expression of specific genes that operate by different mechanisms. These different types of inhibitory polynucleotide technology are described in C. Helene and J. Toulme, (1990) *Biochim. Biophys. Acta.*, 1049:99-125.

[0125] Antisense polynucleotides can include deoxyribonucleotides or ribonucleotides. They can be chemically modified so as to improve stability in the body. Properties of the polynucleotide can be engineered to impart stability (e.g., nuclease resistance), tighter binding or the desired T_m . See, e.g., International patent publication No. 94/12633.

[0126] The general approach to constructing various polynucleotides useful in inhibitory polynucleotide therapy has been reviewed by A.R. Vander Krol *et al.* (1988), *Biotechniques* 6:958-976, and by C.A. Stein *et al.*, (1988) *Cancer Res.* (1988) 48:2659-2668. See also *Oligodeoxynucleotides: Antisense Inhibitors of Gene Expression*, Cohen, J.S., editor, MacMillan Press, London, pages 79-196 (1989), and *Antisense RNA and DNA*, (1988), D.A. Melton, Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY. In certain embodiments inhibitory polynucleotides comprise a derivatized substituent which is substantially non-interfering with respect to hybridization of the inhibitory polynucleotide to the target polynucleotide.

B. Antisense

[0127] This invention provides antisense polynucleotides capable of specifically hybridizing to a target sequence of Repro-EN-1.0 or IB1. Antisense polynucleotides are useful *in vitro* or *in vivo* to inhibit the activity of Repro-EN-1.0 or IB1.

[0128] The antisense polynucleotides of this invention comprise an antisense sequence of at least 7 nucleotides that specifically hybridize to a sequence from Repro-EN-1.0 or IB1 and, more particularly, mammalian Repro-EN-1.0 or IB1 and human Repro-EN-1.0 or IB1.

[0129] The antisense sequence can be between about 10 and about 50 nucleotides or between about 15 and about 35 nucleotides. In other embodiments, antisense polynucleotides are polynucleotides of less than about 100 nucleotides or less than about 200 nucleotides. Accordingly, a sequence of the antisense polynucleotide can specifically

hybridize to all or part of the Repro-EN-1.0 or IB1, such as antisense polynucleotides to the Repro-EN-1.0 or IB1 gene or its transcribed RNA. In one embodiment, the sequence of the polynucleotide contains within it the antisense sequence. In this case, the antisense sequence is contained within a polynucleotide of longer sequence. In another embodiment, the sequence of the polynucleotide consists essentially of, or is, the antisense sequence. Thus, for example, the antisense polynucleotide can be a polynucleotide of less than about 50 nucleotides in a sequence that specifically hybridizes to the target sequence.

[0130] Generally, to assure specific hybridization, the antisense sequence is substantially complementary to the target sequence in Repro-EN-1.0 or IB1. In certain embodiments, the antisense sequence is exactly complementary to the target sequence. The antisense polynucleotides may include nucleotide substitutions, additions, deletions, or transpositions, so long as specific binding to the relevant target sequence corresponding to Repro-EN-1.0 mRNA or its gene, or IB1 mRNA or its gene, is retained as a functional property of the polynucleotide.

[0131] The antisense polynucleotide should be long enough to form a stable duplex but short enough, depending on the mode of delivery, to administer *in vivo*, if desired. The minimum length of a polynucleotide required for specific hybridization to a target sequence depends on several factors, such as G/C content, positioning of mismatched bases (if any), degree of uniqueness of the sequence as compared to the population of target polynucleotides, and chemical nature of the polynucleotide (e.g., methylphosphonate backbone, polyamide nucleic acid, phosphorothioate, etc.), among others.

[0132] For general methods relating to antisense polynucleotides, see *Antisense RNA and DNA*, (1988), D.A. Melton, Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY). For a review of antisense therapy, see, e.g., Uhlmann et al., *Chem. Reviews*, 90:543-584 (1990).

C. Ribozymes

[0133] Cleavage of Repro-EN-1.0 or IB1 can be induced by the use of ribozymes or catalytic RNA. In this approach, the ribozyme would contain either naturally occurring RNA (ribozymes) or synthetic nucleic acids with catalytic activity. Bratty et al., (1992) *Biochim. Biophys. Acta.*, 1216:345-59 (1993) and Denhardt, (1992) *Ann. N. Y. Acad. Sci.*, 660: 70-76 describe methods for making ribozymes.

[0134] Unlike the antisense and other polynucleotides described above, which bind to an RNA or a DNA, a ribozyme not only binds but also specifically cleaves and thereby potentially inactivates a target RNA. Such a ribozyme can comprise 5'- and 3'-terminal sequences complementary to the Repro-EN-1.0 RNA or IB1 RNA.

[0135] Optimum target sites for ribozyme-mediated inhibition of activity can be determined as described by Sullivan et al., PCT patent publication No. 94/02595 and Draper et al., PCT patent publication No. 93/23569. As described by Hu et al., PCT patent publication No. 94/03596, antisense and ribozyme functions can be combined in a single polynucleotide. Upon review of the RNA sequence of Repro-EN-1.0 and IB1, those in the art will note that several useful ribozyme target sites are present and susceptible to cleavage by, for example, a hammerhead motif ribozyme.

[0136] Such engineered ribozymes can be expressed in cells or can be transferred by a variety of means (e.g., liposomes, immunoliposomes, biolistics, direct uptake into cells, etc.). Other forms of ribozymes (group I intron ribozymes (Cech (1995) *Biotechnology* 13; 323); hammerhead ribozymes (Edgington (1992) *Biotechnology* 10: 256) can be engineered on the basis of the disclosed Repro-EN-1.0 or IB1 sequence information to catalyze cleavage of Repro-EN-1.0 RNA or IB1 RNA. Moreover, ribozymes can comprise one or more modified nucleotides or modified linkages between nucleotides, as described above.

D. Other Inhibitory Polynucleotides

[0137] In addition to the antisense and ribozyme inhibitory polynucleotides, one can construct polynucleotides that will bind to duplex nucleic acid either in the folded RNA component or in the gene for the RNA component, forming a triple helix-containing or triplex nucleic acid to inhibit Repro-EN-1.0 or IB1 activity. Such polynucleotides of the invention are constructed using the base-pairing rules of triple helix formation and the nucleotide sequence of the RNA component (Cheng et al. (1988) *J. Biol. Chem.* 263: 15110; Ferrin and Camerini-Otero (1991) *Science* 354: 1494; Ramdas et al. (1989) *J. Biol. Chem.* 264: 17395; Strobel et al. (1991) *Science* 254: 1639; Hsieh et al. (1990) op.cit.; Rigas et al. (1986) *Proc. Natl. Acad. Sci. (U.S.A.)* 83: 9591. Such polynucleotides can block Repro-EN-1.0 or IB1 activity in a number of ways, including by preventing transcription of the Repro-EN-1.0 or IB1 gene.

[0138] Typically, and depending on mode of action, the triplex-forming polynucleotides of the invention comprise a sequence large enough to form a stable triple helix but small enough, depending on the mode of delivery, to administer *in vivo*.

E. Methods for Making Inhibitory Polynucleotides

[0139] Inhibitory polynucleotides can be made chemically or recombinantly.

5 **1. Chemical synthesis**

[0140] Small inhibitory polynucleotides for direct delivery can be made by chemical synthesis. Chemically synthesized polynucleotides can be DNA or RNA, or can include nucleotide analogs or backbones that are not limited to phosphodiester linkages.

10 **2. Recombinant production**

[0141] For delivery into cells or for gene therapy methods, recombinant production of inhibitory polynucleotides through the use of expression vectors is particularly useful. Accordingly, this invention also provides expression vectors, e.g., recombinant polynucleotide molecules comprising expression control sequences operatively linked to the nucleotide sequence encoding the inhibitory polynucleotide.

VII. REPRO-EN-1.0 AND IB1 POLYPEPTIDES

20 [0142] This invention also provides purified, recombinant Repro-EN-1.0 and IB1 polypeptide, and Repro-EN-1.0 and IB1 analogs. Recombinant Repro-EN-1.0 polypeptide includes the polypeptide whose amino acid sequence is presented in SEQ ID NO:2, as well as allelic variants of it. Repro-EN-1.0 analogs include 1) immunogenic fragments of Repro-EN-1.0; 2) homologs of Repro-EN-1.0 from other mammals (especially primates); 3) fragments of Repro-EN-1.0 comprising at least 5 consecutive amino acids from the sequence of Repro-EN-1.0; 4) non-naturally occurring polypeptides whose sequences are substantially identical to Repro-EN-1.0; and 5) fusion proteins comprising a Repro-EN-1.0 or a Repro-EN-1.0 analog fused to a second polypeptide moiety.

[0143] Repro-EN-1.0 polypeptide refers to native Repro-EN-1.0, the polypeptide whose amino acid sequence is the amino acid sequence of SEQ ID NO:2, and to allelic variants of it. Polynucleotide molecules that encode allelic variants of Repro-EN-1.0 are isolatable from endometrial cancer cell cDNA or genomic DNA and typically hybridize under stringent conditions to the nucleotide sequence encoding Repro-EN-1.0 (SEQ ID NO: 1). They can be obtained by amplification using, e.g., PCR primers taken from the sequence of Repro-EN-1.0 described herein.

[0144] Repro-EN-1.0 polypeptides are useful as immunogens to elicit the production of anti-Repro-EN-1.0 antibodies, as affinity capture molecules to isolate such antibodies from a mixture, and as controls in diagnostic methods aimed at detecting Repro-EN-1.0 in a sample.

[0145] Immunogenic Repro-EN-1.0 analogs are polypeptides having a sequence of at least 5 amino acids selected from native Repro-EN-1.0 and which, when presented to an animal as an immunogen, elicit a humoral or cell-mediated immune response. This includes polypeptides comprising an amino acid sequence which is an epitope from Repro-EN-1.0, such as immunogenic fragments of Repro-EN-1.0. Repro-EN-1.0 protein analogs optionally are in isolated form. Persons skilled in the art are familiar with methods of identifying probable epitopes of a protein. For example, polypeptide fragments most likely to elicit an immune response against the native protein are those that exist on the surface of the native protein. Portions of a protein on the surface tend to include hydrophilic amino acids. Such regions can be identified by inspection or with available software. In one embodiment immunogenic polypeptide is a polypeptide comprising a sequence of at least 5 consecutive hydrophilic amino acids, or at least five hydrophilic amino acids in a series of eight consecutive amino acids. Repro-EN-1.0 contains long hydrophilic stretches. Examples of such polypeptides include those comprising the sequences: KTPSAEERR (SEQ ID NO:7), RARPESER (SEQ ID NO:8), RMSDMLSR (SEQ ID NO:9) or NEKLSPKPG (SEQ ID NO:10).

[0146] The cDNA encoding Repro-EN-1.0 of SEQ ID NO:1 was discovered by screening an expression library of cDNA from an endometrial carcinoma cell line with serum pooled from subjects diagnosed with endometriosis. Therefore, polypeptides comprising an epitope of Repro-EN-1.0 can be identified by screening with such serum. Preferably, the test serum is a serum pooled from several subjects positively diagnosed with endometriosis. At least one epitope of Repro-EN-1.0 exists in a fragment of 567 amino acids from the carboxy-terminus of the molecule (amino acids 293-860 of SEQ ID NO:2). At least one epitope of IB1 exists in a fragment of 644 amino acids from the carboxy terminus of the molecule (amino acids 293-937 of SEQ ID NO: 4). Immunogenic fragments are useful, for example, to detect the presence of antibodies against Repro-EN-1.0 or IB1 in patient serum samples. This test is useful in diagnosis because the presence of such antibodies is a diagnostic marker for endometriosis.

[0147] Fragments of Repro-EN-1.0 or IB1 include those having at least 5 amino acids, at least 10 amino acids, at least 50 amino acids, at least 100 amino acids or at least 200 amino acids in a sequence from Repro-EN-1.0 or IB1. Fragments are useful as immunogens to produce an immune response against Repro-EN-1.0 or IB1 in the production

of antibodies. Alternatively, fragments having appropriate amino acid motifs are useful as agretopes to bind with MHC molecules. Such complexes are useful in inducing anergy against Repro-EN-1.0 or IB1.

[0148] Non-naturally occurring analogs of Repro-EN-1.0 or IB1 have at least 90% sequence identity with Repro-EN-1.0 or IB1. They can be made by, for example, introducing conservative amino acid substitutions into the sequence of Repro-EN-1.0 or IB1. Such molecules are useful as decoys or as active analogs.

[0149] Homologs of Repro-EN-1.0 or IB1 from other animals generally have sequences that are substantially identical to that of SEQ ID NO:1 or SEQ ID NO:3. Genomic DNA or cDNA encoding them can be identified through screening libraries under stringent hybridization conditions using a Repro-EN-1.0 or IB1 probe of this invention.

[0150] Fusion proteins include a fragment of Repro-EN-1.0 or IB1 fused to a second polypeptide moiety at the carboxy or amino terminus. The second polypeptide can function, for example, as a detectable label. Such markers include fluorescent protein, enzyme marker of protein from a two-hybrid system.

[0151] Repro-EN-1.0 or IB1 and analogs are most easily produced recombinantly, as described herein. Recombinant Repro-EN-1.0 or IB1 can be purified by affinity purification. In one method, recombinant Repro-EN-1.0 or IB1 analogs comprise a polyhistidine tag. The protein is purified on a nickel-chelate affinity matrix. In another method, Repro-EN-1.0 or IB1 is purified using an affinity matrix carrying anti-Repro-EN-1.0 or IB1 antibodies.

VIII. ANTIBODIES AND HYBRIDOMAS

[0152] In one aspect this invention provides a composition comprising an antibody that specifically binds Repro-EN-1.0 or IB1 polypeptides. Antibodies preferably have affinity of at least 10^6 M⁻¹, 10^7 M⁻¹, 10^8 M⁻¹, or 10^9 M⁻¹. This invention contemplates both polyclonal and monoclonal antibody compositions.

[0153] In one embodiment this invention provides immunotoxins against Repro-EN-1.0- or IB1-expressing cells. Immunotoxins are antibodies and the like as described herein coupled to a compound, e.g., a toxin, that is toxic to a target cell. Toxins can include, for example, radioactive isotopes, ricin, cisplatin, antisense molecules, *Diphtheria* toxin, *Pseudomonas* exotoxin A or *Bacillus anthracis* protective antigen. Immunotoxins bind to cancer cells that express Repro-EN-1.0 or IB1 and kill them. They are useful in the therapeutic methods of this invention. The antibodies of the invention have many uses. For example, such antibodies are useful for detecting Repro-EN-1.0 or IB1 polypeptides in immunoassays. The antibodies also can be used to screen expression libraries for particular expression products such as mammalian Repro-EN-1.0 or IB1. These are useful for detecting or diagnosing various pathological conditions related to the presence of the respective antigens. Usually the antibodies in such a procedure are labeled with a moiety allowing easy detection of presence of antigen by antibody binding. Antibodies raised against Repro-EN-1.0 or IB1 can also be used to raise anti-idiotypic antibodies.

A. Production of Antibodies

[0154] A number of immunogens are used to produce antibodies that specifically bind Repro-EN-1.0 or IB1 polypeptides. Full-length Repro-EN-1.0 or IB1 is a suitable immunogen. Typically, the immunogen of interest is a peptide of at least about 3 amino acids, more typically the peptide is 5 amino acids in length, preferably, the fragment is 10 amino acids in length and more preferably the fragment is 15 amino acids in length or greater. The peptides can be coupled to a carrier protein (e.g., as a fusion protein), or are recombinantly expressed in an immunization vector. Antigenic determinants on peptides to which antibodies bind are typically 3 to 10 amino acids in length. Naturally occurring polypeptides are also used either in pure or impure form.

[0155] Recombinant polypeptides are expressed in eukaryotic or prokaryotic cells and purified using standard techniques. The polypeptide, or a synthetic version thereof, is then injected into an animal capable of producing antibodies. Either monoclonal or polyclonal antibodies can be generated for subsequent use in immunoassays to measure the presence and quantity of the polypeptide.

[0156] Methods for producing polyclonal antibodies are known to those of skill in the art. In brief, an immunogen, preferably a purified polypeptide, a polypeptide coupled to an appropriate carrier (e.g., GST, keyhole limpet hemocyanin, etc.), or a polypeptide incorporated into an immunization vector such as a recombinant vaccinia virus (see, U.S. Patent No. 4,722,848) is mixed with an adjuvant and animals are immunized with the mixture. The animal's immune response to the immunogen preparation is monitored by taking test bleeds and determining the titer of reactivity to the polypeptide of interest. When appropriately high titers of antibody to the immunogen are obtained, blood is collected from the animal and antisera are prepared. Further fractionation of the antisera to enrich for antibodies reactive to the polypeptide is performed where desired. See, e.g., Coligan (1991) *Current Protocols in Immunology* Wiley/Greene, NY; and Harlow and Lane (1989) *Antibodies: A Laboratory Manual* Cold Spring Harbor Press, NY.

[0157] Antibodies, including binding fragments and single chain recombinant versions thereof, against predetermined fragments of Repro-EN-1.0 or IB1 proteins are raised by immunizing animals, e.g., with conjugates of the fragments with carrier proteins as described above.

- [0158] Monoclonal antibodies are prepared from cells secreting the desired antibody. These antibodies are screened for binding to normal or modified polypeptides, or screened for agonistic or antagonistic activity, e.g., activity mediated through Repro-EN-1.0 or IB1. In some instances, it is desirable to prepare monoclonal antibodies from various mammalian hosts, such as mice, rodents, primates, humans, etc. Description of techniques for preparing such monoclonal antibodies are found in, e.g., Stites et al. (eds.) *Basic and Clinical Immunology* (4th ed.) Lange Medical Publications, Los Altos, CA, and references cited therein; Harlow and Lane, *Supra*; Goding (1986) *Monoclonal Antibodies: Principles and Practice* (2d ed.) Academic Press, New York, NY; and Kohler and Milstein (1975) *Nature* 256: 495-497.
- [0159] Other suitable techniques involve selection of libraries of recombinant antibodies in phage or similar vectors. See, Huse et al. (1989) *Science* 246: 1275-1281; and Ward, et al. (1989) *Nature* 341: 544-546.
- [0160] Also, recombinant immunoglobulins may be produced. See, Cabilly, U.S. Patent No. 4,816,567; and Queen et al. (1989) *Proc. Nat'l Acad. Sci. USA* 86: 10029-10033.
- [0161] Frequently, the polypeptides and antibodies will be labeled by joining, either covalently or non-covalently, a substance which provides for a detectable signal. A wide variety of labels and conjugation techniques are known and are reported extensively in both the scientific and patent literature. Thus, an antibody used for detecting an analyte can be directly labeled with a detectable moiety, or may be indirectly labeled by, for example, binding to the antibody a secondary antibody that is, itself directly or indirectly labeled.
- [0162] The antibodies of this invention are also used for affinity chromatography in isolating Repro-EN-1.0 or IB1 proteins. Columns are prepared, e.g., with the antibodies linked to a solid support, e.g., particles, such as agarose, Sephadex, or the like, where a cell lysate is passed through the column, washed, and treated with increasing concentrations of a mild denaturant, whereby purified Repro-EN-1.0 or IB1 polypeptides are released.
- [0163] An alternative approach is the generation of humanized immunoglobulins by linking the CDR regions of non-human antibodies to human constant regions by recombinant DNA techniques. See Queen et al., United States patent 5,585,089.
- [0164] A further approach for isolating DNA sequences which encode a human monoclonal antibody or a binding fragment thereof is by screening a DNA library from human B cells according to the general protocol outlined by Huse et al., *Science* 246:1275-1281 (1989) and then cloning and amplifying the sequences which encode the antibody (or binding fragment) of the desired specificity. The protocol described by Huse is rendered more efficient in combination with phage display technology. See, e.g., Dower et al., WO 91/17271 and McCafferty et al., WO 92/01047. Phage display technology can also be used to mutagenize CDR regions of antibodies previously shown to have affinity for Repro-EN-1.0 or IB1 protein receptors or their ligands. Antibodies having improved binding affinity are selected.
- [0165] In another embodiment of the invention, fragments of antibodies against Repro-EN-1.0 or IB1 protein or protein analogs are provided. Typically, these fragments exhibit specific binding to the Repro-EN-1.0 protein receptor similar to that of a complete immunoglobulin. Antibody fragments include separate heavy chains, light chains Fab, Fab' F(ab')₂ and Fv. Fragments are produced by recombinant DNA techniques, or by enzymic or chemical separation of intact immunoglobulins.

IX. METHODS FOR DETECTING REPRO-EN-1.0 AND IB1 POLYPEPTIDES

- [0166] Repro-EN-1.0 or IB1 polypeptides can be identified by any methods known in the art. In one embodiment, the methods involve detecting the polypeptide with a ligand that specifically recognizes the polypeptide (e.g., an immunoassay). The antibodies of the invention are particularly useful for specific detection of Repro-EN-1.0 or IB1 polypeptides. A variety of antibody-based detection methods are known in the art. These include, for example, radioimmunoassay, sandwich immunoassays (including ELISA), immunofluorescent assays, western blot, affinity chromatography (affinity ligand bound to a solid phase), and *in situ* detection with labeled antibodies. Another method for detecting Repro-EN-1.0 or IB1 polypeptides involves identifying the polypeptide according to its mass through, for example, gel electrophoresis, mass spectrometry or HPLC. Subject samples can be taken from any number of appropriate sources, such as saliva, peritoneal fluid, blood or a blood product (e.g., serum), urine, tissue biopsy (e.g., lymph node tissue), etc.

50 a. Immunoassays

- [0167] The present invention also provides methods for detection of Repro-EN-1.0 or IB1 1 polypeptides employing one or more anti-Repro-EN-1.0 or IB1 1 antibody reagents (i.e., immunoassays). A number of well established immunological binding assay formats suitable for the practice of the invention are known (see, e.g., U.S. Patents 4,366,241; 4,376,110; 4,517,288; and 4,837,168). See, e.g., METHODS IN CELL BIOLOGY VOLUME 37: ANTIBODIES IN CELL BIOLOGY, Asai, ed. Academic Press, Inc. New York (1993); BASIC AND CLINICAL IMMUNOLOGY 7th Edition, Stites & Terr, eds. (1991); Harlow and Lane, *supra* [e.g., Chapter 14], and Ausubel et al., *supra*, [e.g., Chapter 11]. Typically, immunological binding assays (or immunoassays) utilize a "capture agent" to specifically bind to and, often, immobilize

the analyte to a solid phase. In one embodiment, the capture agent is a moiety that specifically binds to a Repro-EN-1.0 or IB1 polypeptide or subsequence, such as an anti-Repro-EN-1.0 or anti-IB1 antibody.

[0168] Usually the Repro-EN-1.0 or IB1 polypeptide being assayed is detected directly or indirectly using a detectable label. The particular label or detectable group used in the assay is usually not a critical aspect of the invention, so long as it does not significantly interfere with the specific binding of the antibody or antibodies used in the assay. The label may be covalently attached to the capture agent (e.g., an anti-Repro-EN-1.0 or anti-IB1 antibody), or may be attached to a third moiety, such as another antibody, that specifically binds to the Repro-EN-1.0 polypeptide.

[0169] The present invention provides methods and reagents for competitive and noncompetitive immunoassays for detecting Repro-EN-1.0 or IB1 polypeptides. Non-competitive immunoassays are assays in which the amount of captured analyte (in this case Repro-EN-1.0 or IB1) is directly measured. One such assay is a two-site, monoclonal-based immunoassay utilizing monoclonal antibodies reactive to two non-interfering epitopes on the Repro-EN-1.0 or IB1 protein. See, e.g., Maddox et al., 1983, *J. Exp. Med.*, 158:1211 for background information. In one preferred "sandwich" assay, the capture agent (e.g., an anti-Repro-EN-1.0 or anti-IB1 antibody) is bound directly to a solid substrate where it is immobilized. These immobilized antibodies then capture any Repro-EN-1.0 or IB1 protein present in the test sample.

[0170] The Repro-EN-1.0 or IB1 polypeptide thus immobilized can then be labeled, i.e., by binding to a second anti-Repro-EN-1.0 or IB1 antibody bearing a label. Alternatively, the second anti-Repro-EN-1.0 or IB1 antibody may lack a label, but be bound by a labeled third antibody specific to antibodies of the species from which the second antibody is derived. The second antibody alternatively can be modified with a detectable moiety, such as biotin, to which a third labeled molecule can specifically bind, such as enzyme-labeled streptavidin.

[0171] In competitive assays, the amount of Repro-EN-1.0 or IB1 protein present in the sample is measured indirectly by measuring the amount of an added (exogenous) Repro-EN-1.0 or IB1 displaced (or competed away) from a capture agent (e.g., anti-Repro-EN-1.0 or anti-IB1 antibody) by the Repro-EN-1.0 or IB1 protein present in the sample.

[0172] A hapten inhibition assay is another example of a competitive assay. In this assay Repro-EN-1.0 or IB1 protein is immobilized on a solid substrate. A known amount of anti-Repro-EN-1.0 or anti-IB1 antibody is added to the sample, and the sample is then contacted with the immobilized Repro-EN-1.0 or IB1 protein. In this case, the amount of anti-Repro-EN-1.0 or anti-IB1 antibody bound to the immobilized Repro-EN-1.0 or IB1 protein is inversely proportional to the amount of Repro-EN-1.0 protein present in the sample. The amount of immobilized antibody may be detected by detecting either the immobilized fraction of antibody or the fraction of the antibody that remains in solution. In this aspect, detection may be direct, where the antibody is labeled, or indirect where the label is bound to a molecule that specifically binds to the antibody as described above.

b. Other Antibody-based Assay Formats

[0173] The invention also provides reagents and methods for detecting and quantifying the presence of Repro-EN-1.0 or IB1 polypeptide in the sample by using an immunoblot (Western blot) format. Another immunoassay is the so-called "lateral flow chromatography." In a non-competitive version of lateral flow chromatography, a sample moves across a substrate by, e.g., capillary action, and encounters a mobile labeled antibody that binds the analyte forming a conjugate. The conjugate then moves across the substrate and encounters an immobilized second antibody that binds the analyte. Thus, immobilized analyte is detected by detecting the labeled antibody. In a competitive version of lateral flow chromatography a labeled version of the analyte moves across the carrier and competes with unlabeled analyte for binding with the immobilized antibody. The greater the amount of the analyte in the sample, the less the binding by labeled analyte and, therefore, the weaker the signal. See, e.g., May et al., U.S. patent 5,622,871 and Rosenstein, U.S. patent 5,591,645.

c. Solid Phases: Substrates, Solid Supports, Membranes, Filters

[0174] As noted *supra*, depending upon the assay, various components, including the antigen, target antibody, or anti-Repro-EN-1.0 or anti-IB1 antibody, may be bound to a solid surface or support (i.e., a substrate, membrane, or filter paper). Many methods for immobilizing biomolecules to a variety of solid surfaces are known in the art. For instance, the solid surface may be a membrane (e.g., nitrocellulose), a microtiter dish (e.g., PVC, polypropylene, or polystyrene), a test tube (glass or plastic), a dipstick (e.g. glass, PVC, polypropylene, polystyrene, latex, and the like), a microcentrifuge tube, or a glass or plastic bead. The desired component may be covalently bound or noncovalently attached through nonspecific bonding.

[0175] A wide variety of organic and inorganic polymers, both natural and synthetic may be employed as the material for the solid surface. Illustrative polymers include polyethylene, polypropylene, poly(4-methylbutene), polystyrene, polymethacrylate, poly(ethylene terephthalate), rayon, nylon, poly(vinyl butyrate), polyvinylidene difluoride (PVDF), silicones, polyformaldehyde, cellulose, cellulose acetate, nitrocellulose, and the like. Other materials which may be employed, include paper, glasses, ceramics, metals, metalloids, semiconductive materials, cements or the like. In

addition, substances that form gels, such as proteins (e.g., gelatins), lipopolysaccharides, silicates, agarose and polyacrylamides can be used. Polymers which form several aqueous phases, such as dextrans, polyalkylene glycols or surfactants, such as phospholipids, long chain (12-24 carbon atoms) alkyl ammonium salts and the like are also suitable. Where the solid surface is porous, various pore sizes may be employed depending upon the nature of the system.

5

d. Mass Spectrometry

[0175] The mass of a molecule frequently can be used as an identifier of the molecule. Therefore, methods of mass spectrometry can be used to identify a protein analyte. Mass spectrometers can measure mass by determining the time required for an ionized analyte to travel down a flight tube and to be detected by an ion detector.

10

[0176] One method of mass spectrometry for proteins is matrix-assisted laser desorption/ionization mass spectrometry ("MALDI"). In MALDI the analyte is mixed with an energy absorbing matrix material that absorbs energy of the wavelength of a laser and placed on the surface of a probe. Upon striking the matrix with the laser, the analyte is desorbed from the probe surface, ionized, and detected by the ion detector. See, for example, Hillenkamp et al., United States patent 5,118,937.

15

[0177] Other methods of mass spectrometry for proteins are described in Hutchens and Yip, United States patent 5,719,060. In one such method referred to as Surfaces Enhanced for Affinity Capture ("SEAC") a solid phase affinity reagent that binds the analyte specifically or non-specifically, such as an antibody or a metal ion, is used to separate the analyte from other materials in a sample. Then the captured analyte is desorbed from the solid phase by, e.g., laser energy, ionized, and detected by the detector.

20

e. Assay Combinations

[0178] The diagnostic and prognostic assays described herein can be carried out in various combinations and can also be carried out in conjunction with other diagnostic or prognostic tests. For example, when the present methods are used to diagnose endometriosis, the presence of a Repro-EN-1.0 or IB1 polypeptide can be used to determine the stage of the disease. Tests that may provide additional information include microscopic analysis of biopsy samples, detection of antigens (e.g., cell-surface markers) associated with endometriosis (e.g., using histocytochemistry, FACS, or the like).

25

X. DIAGNOSTIC, MONITORING AND PROGNOSTIC METHODS

A. Methods Of Diagnosing Endometriosis

[0179] We have detected circulating antibodies against Repro-EN-1.0 or IB1 in the blood of women diagnosed with endometriosis. This supports the idea that endometriosis has an autoimmune component. Further, Repro-EN-1.0 or IB1 and auto-antibodies against Repro-EN-1.0 or IB1 represent two targets in the diagnosis of endometriosis.

30

[0180] Repro-EN-1.0 or IB1 that is shed into the peritoneal fluid of women with endometriosis is useful in methods of diagnosing endometriosis. These methods include detecting Repro-EN-1.0 or IB1 in a biological sample of a subject.

35

Suitable samples include, without limitation, saliva, blood or a blood product (e.g., serum), urine, menstrual fluid, vaginal secretion and, in particular, peritoneal fluid. Repro-EN-1.0 or IB1 can be detected by any of the methods described herein. Any detection of Repro-EN-1.0 or IB1 above a normal range is a positive sign in the diagnosis of endometriosis:

[0181] In another aspect, this invention provides methods for diagnosing endometriosis in a subject by detecting in a sample from the subject a diagnostic amount of an antibody that specifically binds to Repro-EN-1.0 or IB1 polypeptide.

40

Suitable patient samples include, without limitation, saliva, blood or a blood product (e.g., serum), peritoneal fluid, urine, menstrual fluid, vaginal secretion. The antibodies can be detected by any of the methods for detecting proteins described herein. However, sandwich type assays are particularly useful. In one version, all antibodies are captured onto a solid phase, for example using protein A, and antibodies specific for Repro-EN-1.0 or IB1 are detected using a directly or indirectly labeled Repro-EN-1.0 or IB1 or polypeptide fragment of it having an epitope of Repro-EN-1.0 or IB1. In another version of the assay, Repro-EN-1.0 or IB1 or an antigenic fragment of it can be used as the capture molecule and captured antibodies can be detected.

45

[0182] While the detection of antibodies, in general, against Repro-EN-1.0 or IB1 is a positive sign of endometriosis, IgE and IgG₄ class antibodies are particularly specific and sensitive for the diagnosis of endometriosis. Therefore, in one embodiment, the diagnostic method involves specifically detecting IgE or IgG₄ antibodies that specifically recognize Repro-EN-1.0 or IB1. Anti-human IgE antibodies and anti-human IgG₄ antibodies can be easily bought or made.

50

55

1. IN VIVO DIAGNOSIS

[0183] In another method of the invention, endometriosis can be diagnosed *in vivo*. The methods involve detecting Repro-EN-1.0 or IB1 in the body, e.g., in the peritoneum. In general, any conventional method for visualizing diagnostic imaging can be used.

[0184] In one method for diagnosing endometriosis, detection is performed by laparoscopy. A ligand specific for Repro-EN-1.0 or IB1 is introduced into the subject at the site of a suspected lesion and binding is detected using the laparoscope. Alternatively, the binding can be detected by, for example, magnetic resonance imaging (MRI) or electron spin resonance (ESR). Usually gamma-emitting and positron-emitting radioisotopes are used for camera imaging and paramagnetic isotopes are used for magnetic resonance imaging. Any amount of binding above background is a positive sign of endometriosis. Persons of skill in the art recognize that not every positive sign results in a definitive diagnosis of a disease.

[0185] Endometriotic lesions can be removed surgically. However, lesions may be tiny, and difficult to identify by eye. This invention takes advantage of Repro-EN-1.0 or IB1 as marker for endometriosis by providing a method to identify and remove endometriotic lesions. The method involves identifying endometriotic lesions *in situ* using a labeled probe directed to Repro-EN-1.0 or IB1. Then, the lesions are removed surgically.

[0186] In the practice of this method, a probe is provided. The probe binds to Repro-EN-1.0 or IB1 and is labeled with a detectable marker that can be detected in a surgical procedure. In particular, the probe can be an antibody that specifically binds Repro-EN-1.0 or IB1. Preferred labels that can be detected during surgery are radioactive labels and fluorescent labels. Radioactive labels can be detected with the use of, e.g., a Geiger counter. Fluorescent labels, such as FITC, can be detected using, e.g., a D-Light-System (Storz, Tuttlingen Germany).

[0187] Surgery can proceed as follows. The labeled probe is introduced into the peritoneum of the patient for a time sufficient for the label to bind to endometriotic lesions. Unbound labeled probe is washed out. Then, endometriotic lesions are identified using a suitable detector. For example, in laparoscopic surgery, a Geiger counter may be introduced through the incision. Radioactive ("hot") spots indicate bound labeled and, therefore, an endometriotic lesion. These lesions are then removed from the patient.

B. Methods Of Diagnosing Cancer

[0188] Repro-EN-1.0 or IB1 is up-regulated in breast cancer cells, uterine cancer cells, and prostate cancer cells. Therefore, Repro-EN-1.0 or IB1 is a marker for these pathologic conditions. Accordingly, the methods described herein for detecting Repro-EN-1.0 or IB1 polynucleotides or Repro-EN-1.0 or IB1 polypeptides in a sample are useful in methods for diagnosing these cancers, monitoring their progress or treatment, and determining patient prognosis. The methods of the present invention allow cancerous conditions to be detected with increased confidence and at an earlier stage, before cells are detected as cancerous based on pathological characteristics. It is, of course, understood by diagnosticians that diagnostic tests are measured by their degree of specificity and sensitivity. Tests which are not perfectly specific or sensitive are, nevertheless, useful in diagnosis because they provide useful information which, in combination with other evidence, can provide a definitive diagnosis or indicate a course of treatment.

[0189] Methods for diagnosis involve determining a diagnostic amount of Repro-EN-1.0 or IB1 (e.g., mRNA, cDNA or polypeptide) in a patient sample and comparing that amount with a normal range (e.g., a control amount) expected to be found in the sample. The samples used to determine the normal range of Repro-EN-1.0 or IB1 can be normal samples from the individual to be tested, or normal samples from other individuals not suffering from the disease condition.

[0190] A variety of patient samples can be used in the methods of the invention. For example, cell extracts, cultured cells, or tissue samples provide convenient samples for use with the methods of the invention. The methods of the invention can use samples either in solution or extracts, for example, with RT-PCR, or samples such as tissue sections for *in situ* methods of detection. Samples can also be obtained from sources such as fine-needle biopsies, e.g., from breast, uterus or prostate; cellular materials; whole cells; tissue and cell extracts; RNA extracted from tissue and cells and histological sections of tissue.

[0191] Methods for monitoring the course of a cancer with which Repro-EN-1.0 or IB1 is associated involve determining the amount of Repro-EN-1.0 or IB1 in a sample at a first and second time. The times can be during routine physical examinations or during a course of treatment for the cancer. As cancer appears and/or progresses, the amount of Repro-EN-1.0 or IB1 in a sample is expected to increase. Regression or cure of the cancer are accompanied by a decrease or elimination of Repro-EN-1.0 or IB1 in a sample.

[0192] The diagnostic and prognostic methods can also be carried out in conjunction with other diagnostic or prognostic tests. In some instances, such combination tests can provide useful information regarding the progression of a disease, although the present methods for testing for Repro-EN-1.0 or IB1 provide much useful information in this regard.

[0193] Another diagnostic method of the invention involves the administration to a subject of a labeled composition that specifically binds to cells bearing Repro-EN-1.0, or IB1 such as labelled antibodies. Then, the localization of the label is determined by any of the known radiologic methods. Any conventional method for visualizing diagnostic imaging can be used. Usually gamma- and positron-emitting radioisotopes are used for camera imaging and paramagnetic isotopes are used for MRI.

5

C. Methods Of Diagnosing Chromosomal Changes

[0194] The Repro-EN-1.0 or IB1 gene is located on chromosome 1. A translocation at this site can result in alteration 10 of Repro-EN-1.0 or IB1 activity, such as activated transcription or changed function. Chromosomal translocations in the vicinity of the Repro-EN-1.0 or IB1 gene can be detected by hybridizing a labeled probe of this invention to a chromosome spread. A translocation, duplication or deletion can be identified by aberrant hybridization patterns compared to normal. Such tests are useful in detecting genetic abnormalities such as familial disposition to breast, uterine 15 or prostate cancer, or early onset of the disease. A method for fluorescent *in situ* hybridization of chromosomes is provided in the Examples.

15

[0195] The present invention also provides for kits for performing the diagnostic and prognostic method of the invention. Such kits include a polynucleotide probe or primer, or an antibody specific for Repro-EN-1.0 or IB1 and instructions to use the reagents to detect Repro-EN-1.0 or IB1 in a patient sample.

20 XI. METHODS FOR INHIBITING REPRO-EN-1.0 OR IB1 EXPRESSION OR ACTIVITY AND OF TREATING CANCER

[0196] Inhibiting Repro-EN-1.0 or IB1 expression or activity in a breast, uterine or prostate cancer cell can alter the rate of growth or aggressiveness of the cancer. Inhibiting Repro-EN-1.0 or IB1 expression or activity is useful *in vivo* 25 in the prophylactic and therapeutic treatment of prostate cancer or other conditions involving Repro-EN-1.0 or IB1 expression. Accordingly, this invention provides methods for inhibiting Repro-EN-1.0 or IB1 expression or activity. The methods involve contacting a prostate cancer cell, *in vitro* or *in vivo*, with an inhibitory polynucleotide, an immunotoxin or another compound that inhibits Repro-EN-1.0 or IB1 expression or activity.

25

A. Delivery of Inhibitory Polynucleotides

30

[0197] This invention contemplates a variety of means for delivering an inhibitory polynucleotide to a subject including, for example, direct uptake of the molecule by a cell from solution, facilitated uptake through lipofection (e.g., liposomes or immunoliposomes), particle-mediated transfection, and intracellular expression from an expression cassette having an expression control sequence operably linked to a nucleotide sequence that encodes the inhibitory 35 polynucleotide. Methods useful for delivery of polynucleotides for therapeutic purposes are described in *Inouye et al.*, U.S. Patent 5,272,065.

35

B. Pharmaceutical Compositions and Treatment

40

[0198] Agents, such as inhibitory polynucleotides, immunotoxins or other compounds that inhibit Repro-EN-1.0 or IB1 expression or activity preferably are delivered in pharmaceutical compositions comprising the agent and a pharmaceutically acceptable carrier. The agent can be administered by any route that gives it access to cells expressing Repro-EN-1.0 or IB1, for example, prostate tumor cells. This includes, for example, aqueous solutions for enteral, parenteral or transmucosal administration, e.g., for intravenous administration, as tonics and administration to mucous 45 or other membranes as, for example, nose or eye drops; solid and other non-aqueous compositions for enteral or transdermal delivery, e.g., as pills, tablets, powders or capsules; transdermal or transmucosal delivery systems for topical administration, and aerosols or mists for delivery by inhalation. One advantage of delivery by a mode that is easy to administer, e.g., enteral or by intravenous or intramuscular injection is that such modes mimic possible modes of delivery should the agent be formulated as a pharmaceutical.

50

[0199] In one embodiment, the pharmaceutical composition is in the form of a unit dose which contains a pharmacologically effective amount of the Repro-EN-1.0 or IB1 inhibitory compound. The unit dose, taken as part of a therapeutic regimen, results in inhibition of growth of prostate cancer cells. Thus, the pharmaceutical compositions of the invention, whatever the form, are administered in a pharmacologically effective amount to the subject.

55

[0200] The amount of the pharmaceutical composition delivered, the mode of administration and the time course of treatment are at the discretion of the treating physician. Prophylactic treatments are indicated for persons at higher than average risk of getting prostate cancer, breast cancer or uterine cancer. For example, persons with elevated PSA, PAP (prostate acid phosphatase) or PSP (prostate specific protein) levels are at increased risk of prostate cancer. Persons who have the BRCA1 or BRCA2 genes are at increased risk of breast cancer.

XII. METHODS OF INHIBITING AN IMMUNE RESPONSE AGAINST REPRO-EN-1.0 OR IB1

[0201] Women with endometriosis exhibit auto-antibodies against Repro-EN-1.0 or IB1. This fact supports the idea that endometriosis involves an auto-immune response. Thus, this invention provides methods useful for inhibiting an immune response against Repro-EN-1.0 or IB1. In one embodiment, the methods include suppressing the immune system in persons with endometriosis. This includes, for example, the administration of immunosuppressive drugs such as anti-histamines, anti-inflammatories such as steroids, or cyclosporin or anti-idiotypic antibodies that recognize auto-antibodies against Repro-EN-1.0 or IB1.

[0202] In one embodiment of the invention, the immune response can be diminished by the eliciting in the subject anti-idiotypic antibodies against auto-antibodies that specifically recognize Repro-EN-1.0 or IB1. Anti-idiotypic antibodies are produced by immunizing the subject with antibodies against Repro-EN-1.0 or IB1. The amount of delivery to elicit an immune response can be about 1 µg to about 500 µg given with one or more booster administrations over about six weeks. Anti-idiotypic antibodies bind to the antigen binding site of the Repro-EN-1.0 or IB1 antibodies, thereby blocking their function.

[0203] In another embodiment of the invention the method involves inducing anergy in T cells involved in a humoral or cell-mediated immune response against Repro-EN-1.0 or IB1. Anergy can be induced by administering to a subject an MHC-peptide complex that comprises an MHC molecule coupled to a peptide epitope from Repro-EN-1.0 or IB1.

[0204] MHC Class I and MHC Class II molecules bind peptides having particular amino acid motifs in binding pockets located at the amino-terminus of the molecules. The MHC Class II molecule is a dimer formed from an alpha and a beta chain. The binding pocket is formed from portions of both chains. However, a single beta chain suffices to bind a peptide having the appropriate amino acid motif.

[0205] MHC-peptide complexes are formed by contacting a peptide having the appropriate motif with an isolated MHC Class II molecule. Alternatively, the complexes can be formed by creating fusion proteins containing both the MHC molecule and the polypeptide epitope.

[0206] Methods of inducing anergy involve administering an isolated MHC-peptide complex to an individual suffering from the auto-immune disease. Isolated complexes are complexes that do not exist anchored onto a cell surface. MHC-peptide complexes and methods of using them to induce anergy are described in, for example, United States patents 5,468,481 (S.D. Sharma et al.) and 5,734,023 (B. Nag et al.).

[0207] MHC Class II molecules bind peptides having particular amino acid motifs well known in the art. HLA-A1 binding motif includes a first conserved residue of T, S or M, a second conserved residue of D or E, and a third conserved residue of Y. Other second conserved residues are A, S or T. The first and second conserved residues are adjacent and are preferably separated from the third conserved residue by 6 to 7 residues. A second motif consists of a first conserved residue of E or D and a second conserved residue of Y where the first and second conserved residues are separated by 5 to 6 residues. The HLA-A3.2 binding motif includes a first conserved residue of L, M, I, V, S, A, T and F at position 2 and a second conserved residue of K, R or Y at the C-terminal end. Other first conserved residues are C, G or D and alternatively E. Other second conserved residues are H or F. The first and second conserved residues are preferably separated by 6 to 7 residues. The HLA-A11 binding motif includes a first conserved residue of T or V at position 2 and a C-terminal conserved residue of K. The first and second conserved residues are preferably separated by 6 or 7 residues. The HLA-A24.1 binding motif includes a first conserved residue of Y, F or W at position 2 and a C-terminal conserved residue of F, I, W, M or L. The first and second conserved residues are preferably separated by 6 to 7 residues.

[0208] This invention also provides a peptide comprising a linear epitope derived from the Repro-EN-1.0 or IB1, which specifically binds to an MHC molecule. In certain embodiments, the peptide has between 8 and 12 amino acids and the linear epitope has a Class I MHC molecule binding motif.

[0209] The following chart provides portions of the amino acid sequence of Repro-EN-1.0 (SEQ ID NO:2). Amino acid numbers are indicated. Bracketed bars over the amino acid sequence indicate vertebrate MHC Class I or MHC Class II binding motifs. Amino acid numbers are indicated. Peptides of about 8-15 amino acids in length that include these motifs, including peptides whose entire amino acid sequence is selected from the sequence of Repro-EN-1.0, bind to MHC molecules and can be used to induce a cell-mediated or humoral immune response against Repro-EN-1.0. Most of the sequence of Repro-EN-1.0 is exposed on the protein surface and is capable of eliciting an immune response.

[0210] This invention also provides a pharmaceutical composition capable of inducing anergy against Repro-EN-1.0 comprising a pharmaceutically acceptable carrier and an effective amount of an MHC-peptide complex of this invention. The complex is capable of inducing anergy in Class I MHC-restricted cytotoxic T-lymphocytes or Class II MHC-restricted immune response against cells expressing Repro-EN-1.0.

[0211] Repro-EN-1.0 includes many amino acid binding motifs for MHC Class I and MHC Class II. These motifs are provided in Table 1. The amino acid sequence numbers are provided and the motifs are indicated with bars.

TABLE I

337	pnvslmqrmsdmlsrwfeeasevaqsngrgrsrp 347 350 358 361 351 354	(SEQ ID NO:11)
386	vpsspdlevsetamevdtpaeqflq 396 399	(SEQ ID NO:12)
475	- - pvlslhystegtttstiklnftdew 483 489	(SEQ ID NO:13)
536	- - etkapeesssedvtkyqegvsapn 546 549	(SEQ ID NO:14)
561	- - enhinitqsdkftakpldsnsger 571 574	(SEQ ID NO:15)
624	- - ntnpepqfqteatgpsaheetstr 634 636	(SEQ ID NO:16)
675	- - - - - - - - - drrsavariqeffrrtkerkemeeldtlniirplvkrmvykgbrnsrtmikeanfwganfv 685 688 700 703 710 714 720 724	(SEQ ID NO:17)
739	- - - - - dcghifiwdrhtaeilmleadnhvvnclqphpfdfpi 749 752 762 765	(SEQ ID NO:18)
776	- - - - - - - - - - - - - lassgidydiwiwspleesrifurkladevitnelmleetrntitvpasfmlrlmasln 786 789 795 798 804 808 829 833 800 803 810 814	(SEQ ID NO:19)
850	- - - sgqenenedee 855 858	(SEQ ID NO:20)

XIII. TRANSGENIC NON-HUMAN ANIMALS

[0212] This invention also provides non-human mammals transgenic for Repro-EN-1.0 and IB1. As used herein, "animal transgenic for Repro-EN-1.0 or IB1" refers to an animal, in particular a mammal, whose germ cells (i.e., oocytes or sperm), at least, comprise a recombinant nucleic acid molecule comprising expression control sequences operatively linked to a nucleic acid sequence encoding Repro-EN-1.0 or IB1. Such animals are useful, for example, as models in

the study of endometriosis, spontaneous abortion and disease pathways.

[0213] In one embodiment, the expression control sequences are not naturally found operatively linked to Repro-EN-1.0. In one embodiment, the recombinant nucleic acid comprises a non-native Repro-EN-1.0 coding sequence, i.e., a Repro-EN-1.0 sequence that the species does not produce in nature. In one embodiment, the Repro-EN-1.0 is a human Repro-EN-1.0. In another embodiment, the expression control sequences are non-native expression control sequences introduced into the germ cells so as to recombine with the naturally occurring gene and control its expression. Particularly useful transgenic mammals of this invention include rabbits and rodents such as mice.

[0214] The transgenic animals of this invention are produced, for example, by introducing the recombinant nucleic acid molecule into a fertilized egg or embryonic stem (ES) cell, typically by microinjection, electroporation, lipofection, particle-mediated gene transfer. The transgenic animals express the heterologous nucleotide sequence in tissues depending upon whether the promoter is inducible by a signal to the cell, or is constitutive. Transgenic animals can be bred with non-transgenic animals to produce transgenic animals with mixed characteristics.

XIV. METHODS FOR SCREENING FOR COMPOUNDS THAT REGULATE EXPRESSION OF REPRO-EN-1.0 OR IB1

[0215] Compounds that regulate the expression of Repro-EN-1.0 and IB1 are candidates as therapeutic agents in the treatment of breast, uterine or prostate cancer. This invention provides methods for determining whether a compound regulates (e.g., activates or inhibits) expression of Repro-EN-1.0 or IB1.

[0216] Methods for determining whether a compound regulates Repro-EN-1.0 or IB1 expression involve administering to a cell or a test animal having an expressible Repro-EN-1.0 or IB1 gene with the compound, and determining whether expression Repro-EN-1.0 or IB1 is altered. In one embodiment, the methods involve administering the compound to a culture comprising the cell or to a test animal that has cells expressing Repro-EN-1.0 or IB1, measuring the amount of the Repro-EN-1.0 or IB1 polynucleotide or polypeptide in a sample from the culture or the animal, and determining whether the measured amount is different than the amount in a sample from the culture or from the animal under control conditions (e.g., to which no compound has been administered). Statistically significant ($p < 0.05$) differences between the amount measured from the test sample and from the control sample are recorded and indicate that the compound alters the amount of Repro-EN-1.0 or IB1 produced by the cell.

[0217] The compound to be tested can be selected from a number of sources. For example, combinatorial libraries of molecules are available for screening experiments. Using such libraries, thousands of molecules can be screened for regulatory activity. In one preferred embodiment, high throughput screening methods involve providing a library containing a large number of potential therapeutic compounds (candidate compounds). Such "combinatorial chemical libraries" are then screened in one or more assays, as described herein, to identify those library members (particular chemical species or subclasses) that display a desired characteristic activity. The compounds thus identified can serve as conventional "lead compounds" or can themselves be used as potential or actual therapeutics.

[0218] Preparation and screening of combinatorial chemical libraries is well known to those of skill in the art. Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Patent 5,010,175, Furka (1991) *Int. J. Pept. Prot. Res.*, 37: 487-493, Houghton et al. (1991) *Nature*, 354: 84-88). Peptide synthesis is by no means the only approach envisioned and intended for use with the present invention. Other chemistries for generating chemical diversity libraries can also be used. Such chemistries include, but are not limited to: peptoids (PCT Publication No WO 91/19735, 26 Dec. 1991), encoded peptides (PCT Publication WO 93/20242, 14 Oct. 1993), random bio-oligomers (PCT Publication WO 92/00091, 9 Jan. 1992), benzodiazepines (U.S. Pat. No. 5,288,514), diversomers such as hydantoins, benzodiazepines and dipeptides (Hobbs et al., (1993) *Proc. Nat. Acad. Sci. USA* 90: 6909-6913), vinyllogous polypeptides (Hagihara et al. (1992) *J. Amer. Chem. Soc.* 114: 6568), nonpeptidal peptidomimetics with a Beta-D-Glucose scaffolding (Hirschmann et al., (1992) *J. Amer. Chem. Soc.* 114: 9217-9218), analogous organic syntheses of small compound libraries (Chen et al. (1994) *J. Amer. Chem. Soc.* 116: 2661), oligocarbamates (Cho, et al., (1993) *Science* 261:1303), and/or peptidyl phosphonates (Campbell et al., (1994) *J. Org. Chem.* 59: 658). See, generally, Gordon et al., (1994) *J. Med. Chem.* 37:1385, nucleic acid libraries, peptide nucleic acid libraries (see, e.g., U. S. Patent 5,539,083) antibody libraries (see, e.g., Vaughn et al. (1996) *Nature Biotechnology*, 14(3): 309-314), and PCT/US96/10287), carbohydrate libraries (see, e.g., Liang et al. (1996) *Science*, 274: 1520-1522, and U.S. Patent 5,593,853), and small organic molecule libraries (see, e.g., benzodiazepines, Baum (1993) C&EN, Jan 18, page 33, isoprenoids U.S. Patent 5,569,588, thiazolidinones and metathiazanones U.S. Patent 5,549,974, pyrrolidines U.S. Patents 5,525,735 and 5,519,134, morpholino compounds U.S. Patent 5,506,337, benzodiazepines 5,288,514, and the like).

[0219] Devices for the preparation of combinatorial libraries are commercially available (see, e.g., 357 MPS, 390 MPS, Advanced Chem Tech, Louisville KY, Symphony, Rainin, Woburn, MA, 433A Applied Biosystems, Foster City, CA, 9050 Plus, Millipore, Bedford, MA).

[0220] In one embodiment this invention provides inhibitory compounds that inhibit expression of Repro-EN-1.0 or IB1 identified or identifiable by the screening methods of this invention.

XV. GENOMICS

[0221] The identification of cognate or polymorphic forms of the Repro-EN-1.0 or IB1 gene and the tracking of those polymorphisms in individuals and families is important in genetic screening. Accordingly, this invention provides methods useful in detecting polymorphic forms of the Repro-EN-1.0 or IB1 gene. The methods involve comparing the identity of a nucleotide or amino acid at a selected position within the sequence of a test Repro-EN-1.0 or IB1 gene with the nucleotide or amino acid at the corresponding position from the sequence of native Repro-EN-1.0 (SEQ ID NO:1) or IB1 (SEQ ID NO:3). The comparison can be carried out by any methods known in the art, including direct sequence comparison by nucleotide sequencing, sequence comparison or determination by hybridization or identification of RFLPs.

[0222] In one embodiment, the method involves nucleotide or amino acid sequencing of the entire test polynucleotide or polypeptide, or a subsequence from it, and comparing that sequence with the sequence of native Repro-EN-1.0 or IB1. In another embodiment, the method involves identifying restriction fragments produced upon restriction enzyme digestion of the test polynucleotide and comparing those fragments with fragments produced by restriction enzyme digestion of native Repro-EN-1.0 or IB1 gene. Restriction fragments from the native gene can be identified by analysis of the sequence to identify restriction sites. Another embodiment involves the use of oligonucleotide arrays. (See, e.g., Fodor et al., United States patent 5,445,934.) The method involves providing an oligonucleotide array comprising a set of oligonucleotide probes that define sequences selected from the native Repro-EN-1.0 or IB1 sequence, generating hybridization data by performing a hybridization reaction between the target polynucleotide molecules and the probes in the set and detecting hybridization between the target molecules and each of the probes in the set and processing the hybridization data to determine nucleotide positions at which the identity of the target molecule differs from that of native Repro-EN-1.0 or IB1. The comparison can be done manually, but is more conveniently done by a programmable, digital computer.

[0223] The following examples are offered by way of illustration, not by way of limitation.

EXAMPLES

I. Construction of a human endometrial carcinoma cell line (RL95-2) cDNA expression library

A. Material and Methods

[0224] Poly A+ RNA isolated from RL95-2, was used as a template for first strand cDNA synthesis. The poly A+ RNA was analyzed by denaturing gel electrophoresis and ranged in size from 0.2 to 10 Kb (10). An oligo dT primer containing an internal, protected *Xho*I site was annealed in the presence of a nucleotide mixture containing 5-methyl dCTP, and extended with MMTV reverse transcriptase.

[0225] Second strand synthesis of the RL95-2 cDNA/RNA hybrid was completed by the addition of RNase H, DNA polymerase 1, and dNTPs to the first strand synthesis reaction. *Pfu* DNA polymerase was used to blunt-end the double stranded RL95-2 cDNA followed by the ligation of *Eco*R*I* adapters. The cDNA was kinased and digested with *Xho*I and *Eco*R*I* before size fractionation on Sephadryl S-500 columns. The size fractionated cDNA was recovered and the quantified on ethidium bromide containing plates against a set of serially-diluted DNA standards. The cDNA contained in the first two column fractionations was directionally ligated, in the sense orientation, to *Xho*I/*Eco*R*I*-digested uniZAP phage vector arms. Initially, approximately 25 ng (per fraction) of the cDNA was ligated and packaged into bacteriophage particles. Subsequently, the approximately 100 ng remaining cDNA in fractions 1 and 2 was packaged into bacteriophage particles using several reactions of the lambda phage packaging extract (Stratagene).

[0226] After packaging, the primary human RL95-2 library was titered by infection of the XL1 Blue host strain. The ratio of recombinant:nonrecombinant phage was determined by plating infected XL1 Blue in the presence of IPTG and XGal. The number of blue (non-recombinant) or white (recombinant) plaques were quantified using a Manostat colony counter. Ninety-eight and one-half (98.5) percent of the phage in the human RL95-2 cDNA library were recombinant and the primary phage library base consisted of 2.2×10^6 independent clones. The human RL95-2 cDNA library was amplified once and re-titered as above. The titer of the amplified library was 3×10^9 pfu/ml.

[0227] The average size of the cDNA inserts was determined by PCR amplification. Twenty well-isolated phage plaques were cored and the cDNA inserts were amplified using T7/T3 specific oligonucleotides which hybridize to sites flanking the cDNA insertion site contained within the lambda phage vector. The PCR products were analyzed by agarose gel electrophoresis and visualized by ethidium bromide staining. Ninety-five percent of the cored plaques contained inserts varying in size from 0.5 to 2.5 Kb with an average size of 1.5 Kb.

II. Library Screening

[0228] To identify endometrial autoantigens that could be used to develop an endometriosis immunoassay, pooled sera from patients (n=17) with laparoscopy confirmed endometriosis was used as a probe to screen the RL95-2 cDNA expression library using the following methods.

A. Preadsorption of serum. Antibodies that react with expression library host strains were immunoabsorbed from patient serum by using BNN97 and Y1090 *E. coli* lysate-conjugated sepharose beads (5' → 3') following the manufacturer's protocols. Briefly, 2 ml of host stain lysate-conjugated sepharose beads were washed twice with sterile Tris buffered saline (TBS). The beads were resuspended in 4 mls of serum diluted 1/2 in TBS. Following a 16 hour incubation at 4°C, the sepharose beads were collected by centrifugation at 1,000 x g for 2 min. The supernatant was removed and the beads were washed with 4 mls of sterile TBS. After centrifugation at 1,000 x g for 2 min, the supernatants were collected, pooled and used to screen the RL95-2-specific cDNA expression library.

B. Screening the RL95-2 cDNA library. Approximately 10⁶ infectious phage particles were incubated with XL-1 blue host cells and plated at density of 50,000 phage per 150 mm dish using standard protocols (Stratagene). After incubating for 5 hours at 42°C, the phage plaques were overlaid with nitrocellulose membranes (Protran; Schleicher & Schuell) that had been soaked in a 10 mM isopropyl-1-thio-β-D-galactopyranoside (IPTG) solution. Following a 4 hour incubation at 37°C, the membranes were removed and washed three times for 15 min in TBS with 0.05% Tween₂₀ (TTBS). The membranes were incubated with blocking solution (1% Bovine serum albumin [fraction V] in TBS) for 1 hour at room temperature prior to a 1 hour incubation with preadsorbed patient serum diluted 1/10 (final dilution of 1/40) in blocking solution. The membranes were washed three time for 15 min with TTBS prior to the addition of alkaline phosphatase-conjugated goat anti-human Ig(G,A,M) (Pierce) diluted 1/25,000 in blocking solution. After a 1 hour incubation at room temperature the membranes were washed three times with TTBS as described above and once with TBS. The membranes were incubated with enzyme substrate (Western Blue; Promega) for approximately 30 min and the enzymatic reaction was terminated by briefly incubating the membranes with stop solution (Tris-HCl pH 2.9; 1 mM EDTA). Several immunoreactive phage plaques were selected and transferred to 500 µl of SM buffer (100 mM NaCl, 8 mM MgSO₄, 50 mM Tris-HCl pH 7.5, 0.01% gelatin) containing 20 µl of chloroform. The selected phage were eluted from the agar and plated at a density of approximately 1,000 phage per 100 mm dish and screen as described above. To insure that the selected phage plaque, named Repro-EN-1.0, represented a single clone the screening process was repeated a third time as described above.

C. Excision of Repro-EN-1.0 phagemid. Plasmid containing the cDNA insert for Repro-EN-1.0 was excised from the phage clone using the manufacturer's protocols (Stratagene). The size of the Repro-EN-1.0 insert was determined by releasing the cDNA fragment from the rescued pBluescript/Repro-EN-1.0 plasmid with the restriction enzymes *EcoRI* and *XbaI*. The released insert was size fractionated by agarose-gel electrophoresis and the apparent length of the insert was determined by comparing its migration position with a DNA standard (1 kb ladder; Gibco BRL). The insert migrated at approximately 2.0 Kb.

D. Identification of the IB1 clone. An alternately spliced variant of Repro-EN-1.0. A commercial human heart cDNA library (Clontech) was screened with a radiolabeled probe mapping within the amino terminus of the Repro-EN-1.0 coding sequence (nt 203 to 897). One of the two clones isolated contained a cDNA insert of 3.4 Kb which possessed an extra 231 base pair insert within the Repro-EN-1.0 coding sequence.

III. Characterization

A. Sequence analysis

[0229] The nucleotide sequence of Repro-EN-1.0 was determined by using a modified protocol of the dideoxy chain termination method of Sanger et al. and USB Sequenase 2.0 (Barker, D.F. 1993, *Biotechniques*). The amino acid sequence was predicted using the Intelligenetic TRANSLATE program and sequence homologies were determined with BLAST data base search algorithms. The deduced amino acid sequence (in the expected frame for a fusion protein) was novel.

B. Tissue expression analysis

[0230] The Repro-EN-1.0 expression distribution was determined by Northern blot analysis using poly A+ RNA col-

lected from human spleen, thymus, prostate, testis, ovary, small intestine, colon, and peripheral blood leukocytes (MTN human blot 11; Clontech) and human heart, brain, placenta, lung, liver, skeletal muscle, kidney and pancreas (MTN human blot 1- Clontech) and the manufacturer' suggested protocols (Clontech). The immobilized poly A+ RNA samples were incubated with a random prime labeled probe that represented the Repro-EN-1.0 insert. The probe was generated by using and *EcoRI/Xhol* released fragment of pRepro-EN-1.0 as template for a random prime labeled reaction as described in the manufacturer's manual (Megaprime kit; Amersham). The Northern blot membranes were prehybridized with 6 mls of ExpressHyb solution (Clontech) followed by a 1 hour incubation at 68°C with approximately 0.5 ng of radiolabeled probe (approx. 2X10⁵ cpm) in 5 mls of ExpressHyb solution. Unbound probe was removed by washing the membranes three times with 2 X SSC, 0.05% SDS at room temperature for 10 min for each wash followed by twice with 0.1 X SSC, 0.1% SDS at 50° C for 15 min for each wash. The apparent length of RNA species, a 3.4 Kb mRNA, and tissue distribution was determined by autoradiography. Expression was detected primarily in skeletal muscle, heart and testis; and to a lesser extent in other tissues, but was not detected in lung or peripheral blood mononuclear cells (PBMC). Expression of Repro-EN-1.0 was up-regulated in breast and uterine carcinomas relative to their normal counterparts, was highly expressed in both normal fallopian tube and fallopian tube carcinoma, and was expressed at low levels in both normal ovary and ovarian carcinoma (Fig. 2 and Fig. 3). Expression of Repro-EN-1.0 in RL95-2 (endometrium carcinoma) cells is lower than in LNCaP (human prostate adenocarcinoma), PC-12 (rat cell line) and BT12 (human breast carcinoma cell line) cells and undetectable in a mouse hybridoma cell line (3E10; negative control) (Fig. 4). In addition, expression in normal endometrium is undetectable.

20 C. Homologue analysis

[0231] To determine the level of nucleotide conservation of Repro-EN-1.0 in different species, a Southern blot analysis using *EcoRI* digested genomic DNA collected from human, monkey, rat, mouse, dog, cow, rabbit, chicken and yeast was performed as described in the manufacture's manual (ZooBlot; Clontech). Briefly, the immobilized *EcoRI* digested genomic DNA samples were prehybridized with 6 mls of ExpressHyb (Clontech) and incubated for 1 hour at 68°C with 1.0 ng (approx. 4X10⁵ cpm in 5 mls) of a random prime labeled probe that represented the Repro-EN-1.0 insert. Unbound probe was removed by washing the membranes once with 2 X SSC, 0.05% SDS at room temperature for 30 min followed by one wash with 0.1 X SSC, 0.1% SDS at 50C for 30 min. Identification of homologues in different species was determine by autoradiography. (See Figure 4) The sequence is highly conserved between human and non-human primates (Monkey).

IV. Antibodies

[0232] Antibodies to peptides of the clone and/or to recombinant protein were generated in rabbits. This antisera was used to develop a Repro-EN-1.0 ELISA.

V. Recombinant Protein

[0233] The predicted ORF for Repro-EN-1 was subcloned into an expression vector, and the recombinant protein was expressed and purified by Ni--Chelate chromatography using standard methodologies.

VI. ELISA

[0234] The purified recombinant Repro-EN-1 protein was used as a target antigen in an ELISA (EndX™ ELISA) designed to detect antigen-specific autoantibodies in patient serum.

[0235] The present invention provides a novel nucleotide sequence encoding Repro-EN-1.0, Repro-EN-1.0 polypeptides, IB1, IB1 polypeptides and methods of using these materials. While specific examples have been provided, the above description is illustrative and not restrictive. Many variations of the invention will become apparent to those skilled in the art upon review of this specification. The scope of the invention should, therefore, be determined not with reference to the above description, but instead should be determined with reference to the appended claims along with their full scope of equivalents.

[0236] All publications and patent documents cited in this application are incorporated by reference in their entirety for all purposes to the same extent as if each individual publication or patent document were so individually denoted. By their citation of various references in this document Applicants do not admit that any particular reference is "prior art" to their invention.

SEQUENCE LISTING

5 <110> Diagnostic Products Corporation

10 <120> Polynucleotide Encoding Autoantigens Associates With Endometriosis

15 <130> N80980

20 <160> 20

25 <170> PatentIn Ver. 2.1

30 <210> 1

35 <211> 3189

40 <212> DNA

45 <213> Homo sapiens

50 <220>

55 <221> CDS

60 <222> (176)..(2755)

65 <400> 1

70 cggccgggct tcaggggccc aggcggccgt gctgccacgg ccatctaacg ctgcgcacctg 60

75 gaggccccggc gcgcggatgg tgccggtgcg gtcgggtgt tgaaacgggt gtccccctccc 120

80 cctccctcccc tcccccacgc ggtggtctcc cctcccaccc ggctcaggca gagcc atg 178

85 Met

1

90 tct cgg ggt ggc tcc tac cca cac ctg ttg tgg gac gtg agg aaa agg 226
Ser Arg Gly Ser Tyr Pro His Leu Leu Trp Asp Val Arg Lys Arg

95 5

10 10

15

105 ttc ctc ggg ctg gag gac ccg tcc egg ctg egg agt cgc tac ctg gga 274
Phe Leu Gly Leu Glu Asp Pro Ser Arg Leu Arg Ser Arg Tyr Leu Gly

50

55

	20	25	30	
5	aga aga gaa ttt atc caa aga tta aaa ctt gaa gca acc ctt aat gtg			322
	Arg Arg Glu Phe Ile Gln Arg Leu Lys Leu Glu Ala Thr Leu Asn Val			
	35	40	45	
10	cat gat ggt tgt gtt aat aca atc tgt tgg aat gac act gga gaa tat			370
	His Asp Gly Cys Val Asn Thr Ile Cys Trp Asn Asp Thr Gly Glu Tyr			
	50	55	60	65
15	att tta tct ggc tca gat gac acc aaa tta gta att agt aat cct tac			418
	Ile Leu Ser Gly Ser Asp Asp Thr Lys Leu Val Ile Ser Asn Pro Tyr			
	70	75	80	
20	agc aga aag gtt ttg aca aca att cgt tca ggg cac cga gca aac ata			466
	Ser Arg Lys Val Leu Thr Thr Ile Arg Ser Gly His Arg Ala Asn Ile			
	85	90	95	
25	ttt agt gca aag ttc tta cct tgt aca aat gat aaa cag att gta tcc			514
	Phe Ser Ala Lys Phe Leu Pro Cys Thr Asn Asp Lys Gln Ile Val Ser			
	100	105	110	
30	tgc tct gga gat gga gta ata ttt tat acc aac gtt gag caa gat gca			562
	Cys Ser Gly Asp Gly Val Ile Phe Tyr Thr Asn Val Glu Gln Asp Ala			
	115	120	125	
35	gaa acc aac aga caa tgc caa ttt acg tgt cat tat gga act act tat			610
	Glu Thr Asn Arg Gln Cys Gln Phe Thr Cys His Tyr Gly Thr Thr Tyr			
	130	135	140	145
40	gag att atg act gta ccc aat gac cct tac act ttt ctc tct tgt ggt			658
	Glu Ile Met Thr Val Pro Asn Asp Pro Tyr Thr Phe Leu Ser Cys Gly			
	150	155	160	
45	gaa gat gga act gtt agg tgg ttt gat aca cgc atc aaa act agc tgc			706
	Glu Asp Gly Thr Val Arg Trp Phe Asp Thr Arg Ile Lys Thr Ser Cys			
	165	170	175	
50	aca aaa gaa gat tgt aaa gat gat att tta att aac tgt cga cgt gct			754
	Thr Lys Glu Asp Cys Lys Asp Asp Ile Leu Ile Asn Cys Arg Arg Ala			
	180	185	190	
55	gcc acg tct gtt gct att tgc cca cca ata cca tat tac ctt gct gtt			802
	Ala Thr Ser Val Ala Ile Cys Pro Pro Ile Pro Tyr Tyr Leu Ala Val			
	195	200	205	
	ggt tgt tct gac agc tca gta cga ata tat gat cgg cga atg ctg ggc			850
	Gly Cys Ser Asp Ser Ser Val Arg Ile Tyr Asp Arg Arg Met Leu Gly			

EP 1106 690 A2

	210	215	220	225	
5	aca aga gct aca ggg aat tat gca ggt cga ggg act act gga atg gtt				898
	Thr Arg Ala Thr Gly Asn Tyr Ala Gly Arg Gly Thr Thr Gly Met Val				
	230	235	240		
10	gcc cgt ttt att cct tcc cat ctt aat aat aag tcc tgc aga gtg aca				946
	Ala Arg Phe Ile Pro Ser His Leu Asn Asn Lys Ser Cys Arg Val Thr				
	245	250	255		
15	tct ctg tgt tac agt gaa gat ggt caa gag att ctc gtt agt tac tct				994
	Ser Leu Cys Tyr Ser Glu Asp Gly Gln Glu Ile Leu Val Ser Tyr Ser				
	260	265	270		
20	tca gat tac ata tat ctt ttt gac ccg aaa gat gat aca gca cga gaa				1042
	Ser Asp Tyr Ile Tyr Lou Phe Asp Pro Lys Asp Asp Thr Ala Arg Glu				
	275	280	285		
25	ctt aaa act cct tct gcg gaa gag aga aga gaa gag ttg cga caa cca				1090
	Leu Lys Thr Pro Ser Ala Glu Glu Arg Arg Glu Glu Leu Arg Gln Pro				
	290	295	300	305	
30	cca gtt aag cgt ttg aga ctt cgt ggt gat tgg tca gat act gga ccc				1138
	Pro Val Lys Arg Leu Arg Leu Arg Gly Asp Trp Ser Asp Thr Gly Pro				
	310	315	320		
35	aga gca agg ccg gag agt gaa cga gaa cga gat gga gag cag agt ccc				1186
	Arg Ala Arg Pro Glu Ser Glu Arg Glu Arg Asp Gly Glu Gln Ser Pro				
	325	330	335		
40	aat gtg tca ttg atg cag aga atg tct gat atg tta tca aga tgg ttt				1234
	Asn Val Ser Leu Met Gln Arg Met Ser Asp Met Leu Ser Arg Trp Phe				
	340	345	350		
45	gaa gaa gca agt gag gtt gca caa agc aat aga gga cga gga aga tct				1282
	Glu Glu Ala Ser Glu Val Ala Gln Ser Asn Arg Gly Arg Gly Arg Ser				
	355	360	365		
50	cga ccc aga ggt gga aca agt caa tca gat att tca act ctt cct acg				1330
	Arg Pro Arg Gly Gly Thr Ser Gln Ser Asp Ile Ser Thr Leu Pro Thr				
	370	375	380	385	
55	gtc cca tca agt cct gat ttg gaa gtg agt gaa act gca atg gaa gta				1378
	Val Pro Ser Ser Pro Asp Leu Glu Val Ser Glu Thr Ala Met Glu Val				
	390	395	400		
60	gat act cca gct gaa caa ttt ctt cag cct tct aca tcc tct aca atg				1426
	Asp Thr Pro Ala Glu Gln Phe Leu Gln Pro Ser Thr Ser Thr Met				

	405	410	415	
5	tca gct cag gct cat tcg aca tca tct ccc aca gaa agc cct cat tct Ser Ala Gln Ala His Ser Thr Ser Ser Pro Thr Glu Ser Pro His Ser			1474
	420	425	430	
10	act cct ttg cta tct tct cca gat agt gaa caa agg cag tct gtt gag Thr Pro Leu Leu Ser Ser Pro Asp Ser Glu Gln Arg Gln Ser Val Glu			1522
	435	440	445	
15	gca tct gga cac cac aca cat cat cag tct gat aac aat aat gaa aag Ala Ser Gly His His Thr His His Gln Ser Asp Asn Asn Asn Glu Lys			1570
	450	455	460	465
20	ctg agc ccc aaa cca ggg aca ggt gaa cca gtt tta agt ttg cac tac Leu Ser Pro Lys Pro Gly Thr Gly Glu Pro Val Leu Ser Leu His Tyr			1618
	470	475	480	
25	agc aca gaa gga aca act aca agc aca ata aaa ctg aac ttt aca gat Ser Thr Glu Gly Thr Thr Ser Thr Ile Lys Leu Asn Phe Thr Asp			1666
	485	490	495	
30	gaa tgg agc agt ata gca tca agt tct aga gga att ggg agc cat tgc Glu Trp Ser Ser Ile Ala Ser Ser Arg Gly Ile Gly Ser His Cys			1714
	500	505	510	
35	aaa tct gag ggt cag gag gaa tct ttc gtc cca cag agc tca gtg caa Lys Ser Glu Gly Gln Glu Ser Phe Val Pro Gln Ser Ser Val Gln			1762
	515	520	525	
40	cca cca gaa gga gac agt gaa aca aaa gct cct gaa gaa tca tca gag Pro Pro Glu Gly Asp Ser Glu Thr Lys Ala Pro Glu Glu Ser Ser Glu			1810
	530	535	540	545
45	gat gtg aca aaa tat cag gaa gga gta tct gca gaa aac cca gtt gag Asp Val Thr Lys Tyr Gln Glu Gly Val Ser Ala Glu Asn Pro Val Glu			1858
	550	555	560	
50	aac cat atc aat ata aca caa tca gat aag ttc aca gcc aag cca ttg Asn His Ile Asn Ile Thr Gln Ser Asp Lys Phe Thr Ala Lys Pro Leu			1906
	565	570	575	
55	gat tcc aac tca gga gaa aga aat gac ctc aat ctt gat cgc tct tgt Asp Ser Asn Ser Gly Glu Arg Asn Asp Leu Asn Leu Asp Arg Ser Cys			1954
	580	585	590	
	ggg gtt cca gaa gaa tct gct tca tct gaa aaa gcc aag gaa cca gaa Gly Val Pro Glu Glu Ser Ala Ser Ser Glu Lys Ala Lys Glu Pro Glu			2002

EP 1 106 690 A2

	595	600	605	
5	act tca gat cag act agc act gag agt gct acc aat gaa aat aac acc			2050
	Thr Ser Asp Gln Thr Ser Thr Glu Ser Ala Thr Asn Glu Asn Asn Thr			
	610	615	620	625
10	aat cct gag cct cag ttc caa aca gaa gcc act ggg cct tca gct cat			2098
	Asn Pro Glu Pro Gln Phe Gln Thr Glu Ala Thr Gly Pro Ser Ala His			
	630	635	640	
15	gaa gaa aca tcc acc agg gac tct gct ctt cag gac aca gat gac agt			2146
	Glu Glu Thr Ser Thr Arg Asp Ser Ala Leu Gln Asp Thr Asp Asp Ser			
	645	650	655	
20	gat gat gac cca gtc ctg atc cca ggt gca agg tat cga gca gga cct			2194
	Asp Asp Asp Pro Val Leu Ile Pro Gly Ala Arg Tyr Arg Ala Gly Pro			
	660	665	670	
25	ggt gat aga cgc tct gct gtt gcc cgt att cag gag ttc ttc aga cgg			2242
	Gly Asp Arg Arg Ser Ala Val Ala Arg Ile Gln Glu Phe Phe Arg Arg			
	675	680	685	
30	aga aaa gaa agg aaa gaa atg gaa ttg gat act ttg aac att aga			2290
	Arg Lys Glu Arg Lys Glu Met Glu Leu Asp Thr Leu Asn Ile Arg			
	690	695	700	705
35	agg ccg cta gta aaa atg gtt tat aaa ggc cat cgc aac tcc agg aca			2338
	Arg Pro Leu Val Lys Met Val Tyr Lys Gly His Arg Asn Ser Arg Thr			
	710	715	720	
40	atg ata aaa gaa gcc aat ttc tgg ggt gct aac ttt gta atg act ggt			2386
	Met Ile Lys Glu Ala Asn Phe Trp Gly Ala Asn Phe Val Met Thr Gly			
	725	730	735	
45	tct gag tgt ggc cac att ttc atc tgg gat cgg cac act gct gag cat			2434
	Ser Glu Cys Gly His Ile Phe Ile Trp Asp Arg His Thr Ala Glu His			
	740	745	750	
50	ttg atg ctt ctg gaa gct gat aat cat gtg gta aac tgc ctg cag cca			2482
	Leu Met Leu Leu Glu Ala Asp Asn His Val Val Asn Cys Leu Gln Pro			
	755	760	765	
55	cat ccg ttt gac cca att tta gcc tca tct ggc ata gat tat gac ata			2530
	His Pro Phe Asp Pro Ile Leu Ala Ser Ser Gly Ile Asp Tyr Asp Ile			
	770	775	780	785
	aag atc tgg tca cca tta gaa gag tca agg att ttt aac cga aaa ctt			2578
	Lys Ile Trp Ser Pro Leu Glu Ser Arg Ile Phe Asn Arg Lys Leu			

	790	795	800		
5	gct gat gaa gtt ata act cga aac gaa ctc atg ctg gaa gaa act aga			2626	
	Ala	Asp	Glu	Val Ile Thr Arg Asn Glu Leu Met Leu Glu Thr Arg	
	805	810	815		
10	aac acc att aca gtt cca gcc tct ttc atg ttg agg atg ttg gct tca			2674	
	Asn	Thr	Ile	Thr Val Pro Ala Ser Phe Met Leu Arg Met Leu Ala Ser	
	820	825	830		
15	ctt aat cat atc cga gct gac cg ^g ttg gag ggt gac aga tca gaa ggc			2722	
	Leu	Asn	His	Ile Arg Ala Asp Arg Leu Glu Gly Asp Arg Ser Glu Gly	
	835	840	845		
20	tct ggt caa gag aat gaa aat gag gat gag gaa taataaactc tttttggcaa			2775	
	Ser	Gly	Gln	Glu Asn Glu Asn Glu Asp Glu Glu	
	850	855	860		
	gcacttaaat gttctgaat ttgtataaga catttattat tttttttctt tacagagat				2835
25	tttagtgcaat ttaaggtaa tggttttgg agttttccc ttttttggg ataacctaac				2895
	atgggtttgg aatgattgtg tgcatttgcatttggagatttggataaaacaa aactagcaga				2955
30	atgtttttaa aacttttgc cgtgtatgag gagtgctaga aaatgcaaaag tgcattttttt				3015
	tccctaacct tcaaatttgg gagcttggat caatgttggaa gaataattttt catcatatgt				3075
	aaaatgttgg tcaaaataaa tttctacact tgccatttgc atgttttgg ctttctaattt				3135
35	aaagaaaactg gttgttttaa gataccctga aaaaaaaaaa aaaaaaaaaa aaaa				3189
40	<210> 2				
	<211> 860				
	<212> PRT				
	<213> Homo sapiens				
45	<400> 2				
	Met Ser Arg Gly Gly Ser Tyr Pro His Leu Leu Trp Asp Val Arg Lys				
	1	5	10	15	
50	Arg Phe Leu Gly Leu Glu Asp Pro Ser Arg Leu Arg Ser Arg Tyr Leu				
	20	25	30		
	Gly Arg Arg Glu Phe Ile Gln Arg Leu Lys Leu Glu Ala Thr Leu Asn				
	35	40	45		

Val His Asp Gly Cys Val Asn Thr Ile Cys Trp Asn Asp Thr Gly Glu
 50 55 60

5 Tyr Ile Leu Ser Gly Ser Asp Asp Thr Lys Leu Val Ile Ser Asn Pro
 65 70 75 80

10 Tyr Ser Arg Lys Val Leu Thr Thr Ile Arg Ser Gly His Arg Ala Asn
 85 90 95

15 Ile Phe Ser Ala Lys Phe Leu Pro Cys Thr Asn Asp Lys Gln Ile Val
 100 105 110

20 Ser Cys Ser Gly Asp Gly Val Ile Phe Tyr Thr Asn Val Glu Gln Asp
 115 120 125

25 Ala Glu Thr Asn Arg Gln Cys Gln Phe Thr Cys His Tyr Gly Thr Thr
 130 135 140

Tyr Glu Ile Met Thr Val Pro Asn Asp Pro Tyr Thr Phe Leu Ser Cys
 145 150 155 160

30 Gly Glu Asp Gly Thr Val Arg Trp Phe Asp Thr Arg Ile Lys Thr Ser
 165 170 175

35 Cys Thr Lys Glu Asp Cys Lys Asp Asp Ile Leu Ile Asn Cys Arg Arg
 180 185 190

Ala Ala Thr Ser Val Ala Ile Cys Pro Pro Ile Pro Tyr Tyr Leu Ala
 195 200 205

40 Val Gly Cys Ser Asp Ser Ser Val Arg Ile Tyr Asp Arg Arg Met Leu
 210 215 220

Gly Thr Arg Ala Thr Gly Asn Tyr Ala Gly Arg Gly Thr Thr Gly Met
 225 230 235 240

45 Val Ala Arg Phe Ile Pro Ser His Leu Asn Asn Lys Ser Cys Arg Val
 245 250 255

Thr Ser Leu Cys Tyr Ser Glu Asp Gly Gln Glu Ile Leu Val Ser Tyr
 260 265 270

50 Ser Ser Asp Tyr Ile Tyr Leu Phe Asp Pro Lys Asp Asp Thr Ala Arg
 275 280 285

55 Glu Leu Lys Thr Pro Ser Ala Glu Glu Arg Arg Glu Glu Leu Arg Gln
 290 295 300

	Pro Pro Val Lys Arg Leu Arg Leu Arg Gly Asp Trp Ser Asp Thr Gly		
305	310	315	320
5	Pro Arg Ala Arg Pro Glu Ser Glu Arg Glu Arg Asp Gly Glu Gln Ser		
	325	330	335
10	Pro Asn Val Ser Leu Met Gln Arg Met Ser Asp Met Leu Ser Arg Trp		
	340	345	350
	Phe Glu Glu Ala Ser Glu Val Ala Gln Ser Asn Arg Gly Arg Gly Arg		
	355	360	365
15	Ser Arg Pro Arg Gly Gly Thr Ser Gln Ser Asp Ile Ser Thr Leu Pro		
	370	375	380
20	Thr Val Pro Ser Ser Pro Asp Leu Glu Val Ser Glu Thr Ala Met Glu		
	385	390	395
	400		
	Val Asp Thr Pro Ala Glu Gln Phe Leu Gln Pro Ser Thr Ser Ser Thr		
	405	410	415
25	Met Ser Ala Gln Ala His Ser Thr Ser Ser Pro Thr Glu Ser Pro His		
	420	425	430
30	Ser Thr Pro Leu Leu Ser Ser Pro Asp Ser Glu Gln Arg Gln Ser Val		
	435	440	445
	Glu Ala Ser Gly His His Thr His His Gln Ser Asp Asn Asn Asn Glu		
	450	455	460
35	Lys Leu Ser Pro Lys Pro Gly Thr Gly Glu Pro Val Leu Ser Leu His		
	465	470	475
	480		
40	Tyr Ser Thr Glu Gly Thr Thr Ser Thr Ile Lys Leu Asn Phe Thr		
	485	490	495
	Asp Glu Trp Ser Ser Ile Ala Ser Ser Arg Gly Ile Gly Ser His		
	500	505	510
45	Cys Lys Ser Glu Gly Gln Glu Glu Ser Phe Val Pro Gln Ser Ser Val		
	515	520	525
50	Gln Pro Pro Glu Gly Asp Ser Glu Thr Lys Ala Pro Glu Glu Ser Ser		
	530	535	540
	Glu Asp Val Thr Lys Tyr Gln Glu Gly Val Ser Ala Glu Asn Pro Val		
	545	550	555
	560		

Glu Asn His Ile Asn Ile Thr Gln Ser Asp Lys Phe Thr Ala Lys Pro
 565 570 575
 5
 Leu Asp Ser Asn Ser Gly Glu Arg Asn Asp Leu Asn Leu Asp Arg Ser
 580 585 590
 10 Cys Gly Val Pro Glu Glu Ser Ala Ser Ser Glu Lys Ala Lys Glu Pro
 595 600 605
 Glu Thr Ser Asp Gln Thr Ser Thr Glu Ser Ala Thr Asn Glu Asn Asn
 610 615 620
 15 Thr Asn Pro Glu Pro Gln Phe Gln Thr Glu Ala Thr Gly Pro Ser Ala
 625 630 635 640
 20 His Glu Glu Thr Ser Thr Arg Asp Ser Ala Leu Gln Asp Thr Asp Asp
 645 650 655
 Ser Asp Asp Asp Pro Val Leu Ile Pro Gly Ala Arg Tyr Arg Ala Gly
 660 665 670
 25 Pro Gly Asp Arg Arg Ser Ala Val Ala Arg Ile Gln Glu Phe Phe Arg
 675 680 685
 30 Arg Arg Lys Glu Arg Lys Glu Met Glu Glu Leu Asp Thr Leu Asn Ile
 690 695 700
 Arg Arg Pro Leu Val Lys Met Val Tyr Lys Gly His Arg Asn Ser Arg
 705 710 715 720
 35 Thr Met Ile Lys Glu Ala Asn Phe Trp Gly Ala Asn Phe Val Met Thr
 725 730 735
 40 Gly Ser Glu Cys Gly His Ile Phe Ile Trp Asp Arg His Thr Ala Glu
 740 745 750
 His Leu Met Leu Leu Glu Ala Asp Asn His Val Val Asn Cys Leu Gln
 755 760 765
 45 Pro His Pro Phe Asp Pro Ile Leu Ala Ser Ser Gly Ile Asp Tyr Asp
 770 775 780
 50 Ile Lys Ile Trp Ser Pro Leu Glu Glu Ser Arg Ile Phe Asn Arg Lys
 785 790 795 800
 Leu Ala Asp Glu Val Ile Thr Arg Asn Glu Leu Met Leu Glu Glu Thr
 805 810 815
 55

Arg Asn Thr Ile Thr Val Pro Ala Ser Phe Met Leu Arg Met Leu Ala
 820 825 830

5 Ser Leu Asn His Ile Arg Ala Asp Arg Leu Glu Gly Asp Arg Ser Glu
 835 840 845

10 Gly Ser Gly Gln Glu Asn Glu Asn Glu Asp Glu Glu
 850 855 860

15 <210> 3
 <211> 3420
 <212> DNA
 <213> Homo sapiens

20 <220>
 <221> CDS
 <222> (176)..(2986)

25 <400> 3
 cggccgggct tcaggggccc aggccggcgt gctgccaccc ccatctaacc ctgcggccctg 60

gaggccccggc gcgcggatgg tgccggtgcg gctcggtgt tgaaacgggt gtccccctccc 120

30 cctccctcccc tcccccacgc ggtggtctcc cctcccaccc ggctcaggca gagcc atg 178
 Met
 1

35 tct cgg ggt ggc tcc tac cca cac ctg ttg tgg gac gtg agg aaa agg 226
 Ser Arg Gly Gly Ser Tyr Pro His Leu Leu Trp Asp Val Arg Lys Arg
 5 10 15

40 ttc ctc ggg ctg gag gac ccg tcc cgg ctg cgg agt cgc tac ctg gga 274
 Phe Leu Gly Leu Glu Asp Pro Ser Arg Leu Arg Ser Arg Tyr Leu Gly
 20 25 30

45 aga aga gaa ttt atc caa aga tta aaa ctt gaa gca acc ctt aat gtg 322
 Arg Arg Glu Phe Ile Gln Arg Leu Lys Leu Glu Ala Thr Leu Asn Val
 35 40 45

50 cat gat ggt tgt gtt aat aca atc tgt tgg aat gac act gga gaa tat 370
 His Asp Gly Cys Val Asn Thr Ile Cys Trp Asn Asp Thr Gly Glu Tyr
 50 55 60 65

55 att tta tct ggc tca gat gac acc aaa tta gta att agt aat cct tac 418
 Ile Leu Ser Gly Ser Asp Asp Thr Lys Leu Val Ile Ser Asn Pro Tyr
 70 75 80

	agc aga aag gtt ttg aca aca att cgt tca ggg cac cga gca aac ata	466
5	Ser Arg Lys Val Leu Thr Thr Ile Arg Ser Gly His Arg Ala Asn Ile	
	85 90 95	
	ttt agt gca aag ttc tta cct tgt aca aat gat aaa cag att gta tcc	514
10	Phe Ser Ala Lys Phe Leu Pro Cys Thr Asn Asp Lys Gln Ile Val Ser	
	100 105 110	
	tgc tct gga gat gga gta ata ttt tat acc aac gtt gag caa gat gca	562
15	Cys Ser Gly Asp Gly Val Ile Phe Tyr Thr Asn Val Glu Gln Asp Ala	
	115 120 125	
	gaa acc aac aga caa tgc caa ttt acg tgt cat tat gga act act tat	610
20	Glu Thr Asn Arg Gln Cys Gln Phe Thr Cys His Tyr Gly Thr Thr Tyr	
	130 135 140 145	
	gag att atg act gta ccc aat gac cct tac act ttt ctc tct tgt ggt	658
25	Glu Ile Met Thr Val Pro Asn Asp Pro Tyr Thr Phe Leu Ser Cys Gly	
	150 155 160	
	gaa gat gga act gtt agg tgg ttt gat aca cgc atc aaa act agc tgc	706
30	Glu Asp Gly Thr Val Arg Trp Phe Asp Thr Arg Ile Lys Thr Ser Cys	
	165 170 175	
	aca aaa gaa gat tgt aaa gat gat att tta att aac tgt cga cgt gct	754
35	Thr Lys Glu Asp Cys Lys Asp Asp Ile Leu Ile Asn Cys Arg Arg Ala	
	180 185 190	
	gcc acg tct gtt gct att tgc cca cca ata cca tat tac ctt gct gtt	802
40	Ala Thr Ser Val Ala Ile Cys Pro Pro Ile Pro Tyr Tyr Leu Ala Val	
	195 200 205	
	ggg tgt tct gac agc tca gta cga ata tat gat cgg cga atg ctg ggc	850
45	Gly Cys Ser Asp Ser Ser Val Arg Ile Tyr Asp Arg Arg Met Leu Gly	
	210 215 220 225	
	aca aga gct aca ggg aat tat gca ggt cga ggg act act gga atg gtt	898
50	Thr Arg Ala Thr Gly Asn Tyr Ala Gly Arg Gly Thr Thr Gly Met Val	
	230 235 240	
	gcc cgt ttt att cct tcc cat ctt aat aat aag tcc tgc aga gtg aca	946
55	Ala Arg Phe Ile Pro Ser His Leu Asn Asn Lys Ser Cys Arg Val Thr	
	245 250 255	
	tct ctg tgt tac agt gaa gat ggt caa gag att ctc gtt agt tac tct	994
60	Ser Leu Cys Tyr Ser Glu Asp Gly Gln Glu Ile Leu Val Ser Tyr Ser	
	260 265 270	

	tca gat tac ata tat ctt ttt gac ccg aaa gat gat aca gca cga gaa	1042
5	Ser Asp Tyr Ile Tyr Leu Phe Asp Pro Lys Asp Asp Thr Ala Arg Glu	
	275 280 285	
	cct aaa act cct tct gcg gaa gag aga aga gaa gag ttg cga caa cca	1090
10	Leu Lys Thr Pro Ser Ala Glu Glu Arg Arg Glu Glu Leu Arg Gln Pro	
	290 295 300 305	
	cca gtt aag cgt ttg aga ctt cgt ggt gat tgg tca gat act gga ccc	1138
15	Pro Val Lys Arg Leu Arg Gly Asp Trp Ser Asp Thr Gly Pro	
	310 315 320	
	aga gca agg ccg gag agt gaa cga gaa cga gat gga gag cag agt ccc	1186
20	Arg Ala Arg Pro Glu Ser Glu Arg Glu Arg Asp Gly Glu Gln Ser Pro	
	325 330 335	
	aat gtg tca ttg atg cag aga atg tct gat atg tta tca aga tgg ttt	1234
	Asn Val Ser Leu Met Gln Arg Met Ser Asp Met Leu Ser Arg Trp Phe	
	340 345 350	
25	gaa gaa gca agt gag gtt gca caa agc aat aga gga cga gga aga tct	1282
	Glu Glu Ala Ser Glu Val Ala Gln Ser Asn Arg Gly Arg Gly Arg Ser	
	355 360 365	
30	cga ccc aga ggt gga aca agt caa tca gat att tca act ctt cct acg	1330
	Arg Pro Arg Gly Gly Thr Ser Gln Ser Asp Ile Ser Thr Leu Pro Thr	
	370 375 380 385	
35	gtc cca tca agt cct gat ttg gaa gtg agt gaa act gca atg gaa gta	1378
	Val Pro Ser Ser Pro Asp Leu Glu Val Ser Glu Thr Ala Met Glu Val	
	390 395 400	
40	gat act cca gct gaa caa ttt ctt cag cct tct aca tcc tct aca atg	1426
	Asp Thr Pro Ala Glu Gln Phe Leu Gln Pro Ser Thr Ser Ser Thr Met	
	405 410 415	
45	tca gct cag gct cat tcg aca tca tct ccc aca gaa agc cct cat tct	1474
	Ser Ala Gln Ala His Ser Thr Ser Ser Pro Thr Glu Ser Pro His Ser	
	420 425 430	
50	act cct ttg cta tct cca gat agt gaa caa agg cag tct gtt gag	1522
	Thr Pro Leu Leu Ser Ser Pro Asp Ser Glu Gln Arg Gln Ser Val Glu	
	435 440 445	
55	gca tct gga cac cac aca cat cat cag tct gaa ttt tta agg ggg cct	1570
	Ala Ser Gly His His His Gln Ser Glu Phe Leu Arg Gly Pro	
	450 455 460 465	

5	gag ata gct ttg ctt cgt aag cgc ctg caa caa ctg agg ctt aag aag Glu Ile Ala Leu Leu Arg Lys Arg Leu Gln Gln Leu Arg Leu Lys Lys 470 475 480	1618
10	gct gag cag cag agg cag caa gag cta gct gca cat acc cag caa cag Ala Glu Gln Gln Arg Gln Glu Leu Ala Ala His Thr Gln Gln Gln 485 490 495	1666
15	cct tcc act tct gat cag tct tct cat gag ggc tct tca cag gac cct Pro Ser Thr Ser Asp Gln Ser Ser His Glu Gly Ser Ser Gln Asp Pro 500 505 510	1714
20	cat gct tca gat tct cct tct gtg gtt aac aaa cag ctc gga tcc His Ala Ser Asp Ser Pro Ser Val Val Asn Lys Gln Leu Gly Ser 515 520 525	1762
25	atg tca ctt gac gag caa cag gat aac aat aat gaa aag ctg agc ccc Met Ser Leu Asp Glu Gln Gln Asp Asn Asn Glu Lys Leu Ser Pro 530 535 540 545	1810
30	aaa cca ggg aca ggt gaa cca gtt tta agt ttg cac tac agc aca gaa Lys Pro Gly Thr Gly Glu Pro Val Leu Ser Leu His Tyr Ser Thr Glu 550 555 560	1858
35	gga aca act aca agc aca ata aaa ctg aac ttt aca gat gaa tgg agc Gly Thr Thr Ser Thr Ile Lys Leu Asn Phe Thr Asp Glu Trp Ser 565 570 575	1906
40	agt ata gca tca agt tct aga gga att ggg agc cat tgc aaa tct gag Ser Ile Ala Ser Ser Arg Gly Ile Gly Ser His Cys Lys Ser Glu 580 585 590	1954
45	ggt cag gag gaa tct ttc gtc cca cag agc tca gtg caa cca cca gaa Gly Gln Glu Glu Ser Phe Val Pro Gln Ser Ser Val Gln Pro Pro Glu 595 600 605	2002
50	gga gac agt gaa aca aaa gct cct gaa gaa tca tca gag gat gtg aca Gly Asp Ser Glu Thr Lys Ala Pro Glu Glu Ser Ser Glu Asp Val Thr 610 615 620 625	2050
55	aaa tat cag gaa gga gta tct gca gaa aac cca gtt gag aac cat atc Lys Tyr Gln Glu Gly Val Ser Ala Glu Asn Pro Val Glu Asn His Ile 630 635 640	2098
	aat ata aca caa tca gat aag ttc aca gcc aag cca ttg gat tcc aac Asn Ile Thr Gln Ser Asp Lys Phe Thr Ala Lys Pro Leu Asp Ser Asn 645 650 655	2146

5	tca gga gaa aga aat gac ctc aat ctt gat cgc tct tgt ggg gtt cca Ser Gly Glu Arg Asn Asp Leu Asn Leu Asp Arg Ser Cys Gly Val Pro 660 665 670	2194
10	gaa gaa tct gct tca tct gaa aaa gcc aag gaa cca gaa act tca gat Glu Glu Ser Ala Ser Ser Glu Lys Ala Lys Glu Pro Glu Thr Ser Asp 675 680 685	2242
15	cag act agc act gag agt gct acc aat gaa aat aac acc aat cct gag Gln Thr Ser Thr Glu Ser Ala Thr Asn Glu Asn Asn Thr Asn Pro Glu 690 695 700 705	2290
20	cct cag ttc caa aca gaa gcc act ggg cct tca gct cat gaa gaa aca Pro Gln Phe Gln Thr Glu Ala Thr Gly Pro Ser Ala His Glu Glu Thr 710 715 720	2338
25	tcc acc agg gac tct gct ctt cag gac aca gat gac agt gat gat gac Ser Thr Arg Asp Ser Ala Leu Gln Asp Thr Asp Asp Ser Asp Asp Asp 725 730 735	2386
30	cca gtc ctg atc cca ggt gca agg tat cga gca gga cct ggt gat aga Pro Val Leu Ile Pro Gly Ala Arg Tyr Arg Ala Gly Pro Gly Asp Arg 740 745 750	2434
35	cgc tct gct gtt gcc cgt att cag gag ttc ttc aga cgg aga aaa gaa Arg Ser Ala Val Ala Arg Ile Gln Glu Phe Phe Arg Arg Arg Lys Glu 755 760 765	2482
40	agg aaa gaa atg gaa gaa ttg gat act ttg aac att aga agg ccg cta Arg Lys Glu Met Glu Glu Leu Asp Thr Leu Asn Ile Arg Arg Pro Leu 770 775 780 785	2530
45	gta aaa atg gtt tat aaa ggc cat cgc aac tcc agg aca atg ata aaa Val Lys Met Val Tyr Lys Gly His Arg Asn Ser Arg Thr Met Ile Lys 790 795 800	2578
50	gaa gcc aat ttc tgg ggt gct aac ttt gta atg act ggt tct gag tgt Glu Ala Asn Phe Trp Gly Ala Asn Phe Val Met Thr Gly Ser Glu Cys 805 810 815	2626
55	ggc cac att ttc atc tgg gat cgg cac act gct gag cat ttg atg ctt Gly His Ile Phe Ile Trp Asp Arg His Thr Ala Glu His Leu Met Leu 820 825 830	2674
	ctg gaa gct gat aat cat gtg gta aac tgc ctg cag cca cat ccg ttt Leu Glu Ala Asp Asn His Val Val Asn Cys Leu Gln Pro His Pro Phe 835 840 845	2722

5 gac cca att tta gcc tca tct ggc ata gat tat gac ata aag atc tgg 2770
 Asp Pro Ile Leu Ala Ser Ser Gly Ile Asp Tyr Asp Ile Lys Ile Trp
 850 855 860 865

10 tca cca tta gaa gag tca agg att ttt aac cga aaa ctt gct gat gaa 2818
 Ser Pro Leu Glu Glu Ser Arg Ile Phe Asn Arg Lys Leu Ala Asp Glu
 870 875 880

15 gtt ata act cga aac gaa ctc atg ctg gaa gaa act aga aac acc att 2866
 Val Ile Thr Arg Asn Glu Leu Met Leu Glu Thr Arg Asn Thr Ile
 885 890 895

20 aca gtt cca gcc tct ttc atg ttg agg atg ttg gct tca ctt aat cat 2914
 Thr Val Pro Ala Ser Phe Met Leu Arg Met Leu Ala Ser Leu Asn His
 900 905 910

25 atc cga gct gac cggttgg ggttgg gac aga tca gaa ggc tct ggt caa 2962
 Ile Arg Ala Asp Arg Leu Glu Gly Asp Arg Ser Glu Gly Ser Gly Gln
 915 920 925

30 gag aat gaa aat gag gat gag gaa taataaaactc tttttggcaa gcacttaaat 3016
 Glu Asn Glu Asn Glu Asp Glu Glu
 930 935

35 gttctgaaat ttgtataaga catttattat tttttttct ttacagagat ttagtgcaat 3076
 tttaaggta tggttttgg agttttccc ttttttggg ataacctaac attggtttgg 3136
 aatgattgtg tgcattgttattt tgggagattt tataaaaacaa aacttagcaga atgtttttaa 3196
 aacttttgc cgtgtatgag gatgtctaga aaatgcaaaag tgcaatattt tccctaaacct 3256

40 tcaaatgtgg gagcttggat caatgttggaa gaataatttt catcatagtg aaaaatgttgg 3316
 ttcaaaataaa ttctacact tgccatttgc atgtttgttgc ctttctaattt aaagaaaactg 3376
 gttgtttaa gataccctga aaaaaaaaaa aaaaaaaaaa aaaa 3420

45 <210> 4
 <211> 937
 <212> PRT
 <213> Homo sapiens

50 <400> 4
 Met Ser Arg Gly Gly Ser Tyr Pro His Leu Leu Trp Asp Val Arg Lys
 55 1 5 10 15

5	Arg Phe Leu Gly Leu Glu Asp Pro Ser Arg Leu Arg Ser Arg Tyr Leu		
	20	25	30
	Gly Arg Arg Glu Phe Ile Gln Arg Leu Lys Leu Glu Ala Thr Leu Asn		
	35	40	45
10	Val His Asp Gly Cys Val Asn Thr Ile Cys Trp Asn Asp Thr Gly Glu		
	50	55	60
15	Tyr Ile Leu Ser Gly Ser Asp Asp Thr Lys Leu Val Ile Ser Asn Pro		
	65	70	75
	Tyr Ser Arg Lys Val Leu Thr Thr Ile Arg Ser Gly His Arg Ala Asn		
	85	90	95
20	Ile Phe Ser Ala Lys Phe Leu Pro Cys Thr Asn Asp Lys Gln Ile Val		
	100	105	110
25	Ser Cys Ser Gly Asp Gly Val Ile Phe Tyr Thr Asn Val Glu Gln Asp		
	115	120	125
	Ala Glu Thr Asn Arg Gln Cys Gln Phe Thr Cys His Tyr Gly Thr Thr		
	130	135	140
30	Tyr Glu Ile Met Thr Val Pro Asn Asp Pro Tyr Thr Phe Leu Ser Cys		
	145	150	155
	160		
35	Gly Glu Asp Gly Thr Val Arg Trp Phe Asp Thr Arg Ile Lys Thr Ser		
	165	170	175
	Cys Thr Lys Glu Asp Cys Lys Asp Asp Ile Leu Ile Asn Cys Arg Arg		
	180	185	190
40	Ala Ala Thr Ser Val Ala Ile Cys Pro Pro Ile Pro Tyr Tyr Leu Ala		
	195	200	205
45	Val Gly Cys Ser Asp Ser Ser Val Arg Ile Tyr Asp Arg Arg Met Leu		
	210	215	220
	Gly Thr Arg Ala Thr Gly Asn Tyr Ala Gly Arg Gly Thr Thr Gly Met		
	225	230	235
	240		
50	Val Ala Arg Phe Ile Pro Ser His Leu Asn Asn Lys Ser Cys Arg Val		
	245	250	255
	Thr Ser Leu Cys Tyr Ser Glu Asp Gly Gln Glu Ile Leu Val Ser Tyr		
	260	265	270

	Ser Ser Asp Tyr Ile Tyr Leu Phe Asp Pro Lys Asp Asp Thr Ala Arg			
5	275	280	285	
	Glu Leu Lys Thr Pro Ser Ala Glu Glu Arg Arg Glu Glu Leu Arg G'n			
10	290	295	300	
	Pro Pro Val Lys Arg Leu Arg Leu Arg Gly Asp Trp Ser Asp Thr Gly			
15	305	310	315	320
	Pro Arg Ala Arg Pro Glu Ser Glu Arg Glu Arg Asp Gly Glu Gln Ser			
20	325	330	335	
	Pro Asn Val Ser Leu Met Gln Arg Met Ser Asp Met Leu Ser Arg Trp			
25	340	345	350	
	Phe Glu Glu Ala S. r Glu Val Ala Gln Ser Asn Arg Gly Arg Gly Arg			
30	355	360	365	
	Ser Arg Pro Arg Gly Gly Thr Ser Gln Ser Asp Ile Ser Thr Leu Pro			
35	370	375	380	
	Thr Val Pro Ser Ser Pro Asp Leu Glu Val Ser Glu Thr Ala Met Glu			
40	385	390	395	400
	Val Asp Thr Pro Ala Glu Gln Phe Leu Gln Pro Ser Thr Ser Thr			
45	405	410	415	
	Met Ser Ala Gln Ala His Ser Thr Ser Ser Pro Thr Glu Ser Pro His			
50	420	425	430	
	Ser Thr Pro Leu Leu Ser Ser Pro Asp Ser Glu Gln Arg Gln Ser Val			
55	435	440	445	
	Glu Ala Ser Gly His His Thr His His Gln Ser Glu Phe Leu Arg Gly			
	450	455	460	
	Pro Glu Ile Ala Leu Leu Arg Lys Arg Leu Gln Gln Leu Arg Leu Lys			
	465	470	475	480
	Lys Ala Glu Gln Gln Arg Gln Gln Glu Leu Ala Ala His Thr Gln Gln			
	485	490	495	
	Gln Pro Ser Thr Ser Asp Gln Ser Ser His Glu Gly Ser Ser Gln Asp			
55	500	505	510	
	Pro His Ala Ser Asp Ser Pro Ser Ser Val Val Asn Lys Gln Leu Gly			
	515	520	525	

5 Ser Met Ser Leu Asp Glu Gln Gln Asp Asn Asn Asn Glu Lys Leu Ser
 530 535 540

10 Pro Lys Pro Gly Thr Gly Glu Pro Val Leu Ser Leu His Tyr Ser Thr
 545 550 555 560

15 Glu Gly Thr Thr Ser Thr Ile Lys Leu Asn Phe Thr Asp Glu Trp
 565 570 575

20 Ser Ser Ile Ala Ser Ser Arg Gly Ile Gly Ser His Cys Lys Ser
 580 585 590

25 Glu Gly Gln Glu Glu Ser Phe Val Pro Gln Ser Ser Val Gln Pro Pro
 595 600 605

30 Glu Gly Asp Ser Glu Thr Lys Ala Pro Glu Glu Ser Ser Glu Asp Val
 610 615 620

35 Thr Lys Tyr Gln Glu Gly Val Ser Ala Glu Asn Pro Val Glu Asn His
 625 630 635 640

40 Ile Asn Ile Thr Gln Ser Asp Lys Phe Thr Ala Lys Pro Leu Asp Ser
 645 650 655

45 Asn Ser Gly Glu Arg Asn Asp Leu Asn Leu Asp Arg Ser Cys Gly Val
 660 665 670

50 Pro Glu Glu Ser Ala Ser Ser Glu Lys Ala Lys Glu Pro Glu Thr Ser
 675 680 685

55 Asp Gln Thr Ser Thr Glu Ser Ala Thr Asn Glu Asn Asn Thr Asn Pro
 690 695 700

60 Glu Pro Gln Phe Gln Thr Glu Ala Thr Gly Pro Ser Ala His Glu Glu
 705 710 715 720

65 Thr Ser Thr Arg Asp Ser Ala Leu Gln Asp Thr Asp Asp Ser Asp Asp
 725 730 735

70 Asp Pro Val Leu Ile Pro Gly Ala Arg Tyr Arg Ala Gly Pro Gly Asp
 740 745 750

75 Arg Arg Ser Ala Val Ala Arg Ile Gln Glu Phe Phe Arg Arg Arg Lys
 755 760 765

80 Glu Arg Lys Glu Met Glu Glu Leu Asp Thr Leu Asn Ile Arg Arg Pro
 770 775 780

5 Leu Val Lys Met Val Tyr Lys Gly His Arg Asn Ser Arg Thr Met Ile
785 790 795 800

10 Lys Glu Ala Asn Phe Trp Gly Ala Asn Phe Val Met Thr Gly Ser G'u
805 810 815

15 Cys Gly His Ile Phe Ile Trp Asp Arg His Thr Ala Glu His Leu Met
820 825 830

20 Leu Leu Glu Ala Asp Asn His Val Val Asn Cys Leu Gln Pro His Pro
835 840 845

25 Phe Asp Pro Ile Leu Ala Ser Ser Gly Ile Asp Tyr Asp Ile Lys Ile
850 855 860

30 Trp Ser Pro Leu Glu Glu Ser Arg Ile Phe Asn Arg Lys Leu Ala Asp
865 870 875 880

35 Glu Val Ile Thr Arg Asn Glu Leu Met Leu Glu Glu Thr Arg Asn Thr
885 890 895

40 Ile Thr Val Pro Ala Ser Phe Met Leu Arg Met Leu Ala Ser Leu Asn
900 905 910

45 His Ile Arg Ala Asp Arg Leu Glu Gly Asp Arg Ser Glu Gly Ser Gly
915 920 925

50 Gln Glu Asn Glu Asn Glu Asp Glu Glu
930 935

40 <210> 5
<211> 21
<212> DNA
<213> Homo sapiens

45 <400> 5
caggacacag atgacagtga t

50 <210> 6
<211> 20
<212> DNA
<213> Homo sapiens

55 <400> 6

20

agagccttct gatctgtcac

5

<210> 7
<211> 9
<212> PRT
<213> Homo sapiens

10

<400> 7
Lys Thr Pro Ser Ala Glu Glu Arg Arg
1 5

15

<210> 8
<211> 8
<212> PRT
<213> Homo sapiens

20

<400> 8
Arg Ala Arg Pro Glu Ser Glu Arg
1 5

25

<210> 9
<211> 7
<212> PRT
<213> Homo sapiens

30

<400> 9
Arg Met Ser Asp Met Ser Arg
1 5

35

<210> 10
<211> 9
<212> PRT
<213> Homo sapiens

40

<400> 10
Asn Glu Lys Leu Ser Pro Lys Pro Gly
1 5

45

<210> 11
<211> 35
<212> PRT
<213> Homo sapiens

50

<400> 11
 Pro Asn Val Ser Leu Met Gln Arg Met Ser Asp Met Leu Ser Arg Trp
 1 5 10 15
 5

Phe Glu Glu Ala Ser Glu Val Ala Gln Ser Asn Arg Gly Arg Gly Arg
 20 25 30
 10 Ser Arg Pro
 35

15 <210> 12
 <211> 25
 <212> PRT
 <213> Homo sapiens

20 <400> 12
 Val Pro Ser Ser Pro Asp Leu Glu Val Ser Glu Thr Ala Met Glu Val
 1 5 10 15
 25 Asp Thr Pro Ala Glu Gln Phe Leu Gln
 20 25

30 <210> 13
 <211> 25
 <212> PRT
 <213> Homo sapiens

35 <400> 13
 Pro Val Leu Ser Leu His Tyr Ser Thr Glu Gly Thr Thr Thr Ser Thr
 1 5 10 15
 40 Ile Lys Leu Asn Phe Thr Asp Glu Trp
 20 25

45 <210> 14
 <211> 24
 <212> PRT
 <213> Homo sapiens

50 <400> 14
 Glu Thr Lys Ala Pro Glu Glu Ser Ser Glu Asp Val Thr Lys Tyr Gln
 1 5 10 15
 55 Glu Gly Val Ser Ala Glu Asn Pro
 20

5 <210> 15
 <211> 25
 <212> PRT
 <213> Homo sapiens

10 <400> 15
 Glu Asn His Ile Asn Ile Thr Gln Ser Asp Lys Phe Thr Ala Lys Pro
 1 5 10 15

15 Leu Asp Ser Asn Ser Gly Glu Arg Asn
 20 25

20 <210> 16
 <211> 24
 <212> PRT
 <213> Homo sapiens

25 <400> 16
 Asn Thr Asn Pro Glu Pro Gln Phe Gln Thr Glu Ala Thr Gly Pro Ser
 1 5 10 15

30 Ala His Glu Glu Thr Ser Thr Arg
 20

35 <210> 17
 <211> 60
 <212> PRT
 <213> Homo sapiens

40 <400> 17
 Asp Arg Arg Ser Ala Val Ala Arg Ile Gln Glu Phe Phe Arg Arg Arg
 1 5 10 15

45 Lys Glu Arg Lys Glu Met Glu Glu Leu Asp Thr Leu Asn Ile Arg Arg
 20 25 30

50 Pro Leu Val Lys Met Val Tyr Lys Gly His Arg Asn Ser Arg Thr Met
 35 40 45

55 Ile Lys Glu Ala Asn Phe Trp Gly Ala Asn Phe Val
 50 55 60

55 <210> 18

<211> 37

<212> PRT

5 <213> Homo sapiens

<400> 18

Glu Cys Gly His Ile Phe Ile Trp Asp Arg His Thr Ala Glu His Leu

10 1 5 10 15

Met Leu Leu Glu Ala Asp Asn His Val Val Asn Cys Leu Gln Pro His

20 20 25 30

15 Pro Phe Asp Pro Ile
35

20 <210> 19

<211> 60

<212> PRT

<213> Homo sapiens

25 <400> 19
Leu Ala Ser Ser Gly Ile Asp Tyr Asp Ile Lys Ile Trp Ser Pro Leu
1 5 10 1530 Glu Glu Ser Arg Ile Phe Asn Arg Lys Leu Ala Asp Glu Val Ile Thr
20 25 30Arg Asn Glu Leu Met Leu Glu Glu Thr Arg Asn Thr Ile Thr Val Pro
35 35 40 4535 Ala Ser Phe Met Leu Arg Met Leu Ala Ser Leu Asn
50 55 60

40 <210> 20

<211> 11

<212> PRT.

45 <213> Homo sapiens

<400> 20

Ser Gly Gln Glu Asn Glu Asn Glu Asp Glu Glu

50 1 5 10

55

Claims

1. A recombinant polynucleotide comprising a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0 (SEQ ID NO: 2), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.
2. The polynucleotide of claim 1 wherein the nucleotide sequence is selected from the Repro-EN-1.0 sequence of SEQ ID NO: 1.
3. The polynucleotide of claim 1 wherein the nucleotide sequence is a native Repro-EN-1.0 nucleotide sequence.
4. The polynucleotide of claim 1 wherein the nucleotide sequence is identical to nucleotides 176 to 2755 of SEQ ID NO: 1.
5. A recombinant polynucleotide comprising a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of IB1 (SEQ ID NO: 4), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.
6. The polynucleotide of claim 5 wherein the nucleotide sequence is selected from the IB 1 sequence of SEQ ID NO: 3.
7. The polynucleotide of claim 5 wherein the nucleotide sequence is a native IB 1 nucleotide sequence.
8. The polynucleotide of claim 5 wherein the nucleotide sequence is identical to nucleotides 176 to 2986 of SEQ ID NO: 3.
9. The polynucleotide of any one of claims 1 to 8 further comprising an expression control sequence operatively linked to the nucleotide sequence.
10. A polynucleotide primer pair which amplifies a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0, wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis, the pair comprising:
 - (1) a 3' primer of at least 7 nucleotides that specifically hybridizes to a 3' end of the nucleotide sequence or downstream from the sequence, and
 - (2) a 5' primer of at least 7 nucleotides that specifically hybridizes to the 3' end of the complement of the nucleotide sequence or downstream from the complement of the sequence.
11. The polynucleotide primer pair of claim 10 wherein the 3' primer has a sequence complementary to a nucleotide sequence selected from Repro-EN-1.0 cDNA (SEQ ID NO: 1), and the 5' primer has a sequence identical to nucleotide sequence selected from Repro-EN-1.0 cDNA (SEQ ID NO: 1).
12. A polynucleotide primer pair which amplifies a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of IB1, wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis, the pair comprising:
 - (1) a 3' primer of at least 7 nucleotides that specifically hybridizes to a 3' end of the nucleotide sequence or downstream from the sequence, and
 - (2) a 5' primer of at least 7 nucleotides that specifically hybridizes to the 3' end of the complement of the nucleotide sequence or downstream from the complement of the sequence.
13. The polynucleotide primer pair of claim 12 wherein the 3' primer has a sequence complementary to a nucleotide sequence selected from IB1 cDNA (SEQ ID NO: 3), and the 5' primer has a sequence identical to nucleotide sequence selected from IB1 cDNA (SEQ ID NO: 3).
14. The polynucleotide primer pair of claim 10 or 12 wherein the pair of polynucleotides are peptide nucleic acids.
15. A recombinant cell comprising a recombinant polynucleotide comprising an expression control sequence operatively linked to a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of Repro-EN-1.0

(SEQ ID NO: 2), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.

16. A method for detecting a target polynucleotide comprising a nucleotide sequence selected from Repro-EN-1.0 cDNA (SEQ ID NO: 1) or its complement in a sample comprising the steps of:

- 5 (a) contacting the sample with a polynucleotide probe or primer comprising a sequence of at least 7 nucleotides that specifically hybridizes to the nucleotide sequence and
(b) detecting whether the probe or primer has specifically hybridized to the target polynucleotide,

10 whereby specific hybridization provides a detection of the target polynucleotide in the sample.

17. A recombinant cell comprising a recombinant polynucleotide comprising an expression control sequence operatively linked to a nucleotide sequence encoding a polypeptide epitope of at least 5 amino acids of IB1 (SEQ ID NO: 4), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.

18. A method for detecting a target polynucleotide comprising a nucleotide sequence selected from IB1 cDNA (SEQ ID NO: 3) or its complement in a sample comprising the steps of:

- 20 (a) contacting the sample with a polynucleotide probe or primer comprising a sequence of at least 7 nucleotides that specifically hybridizes to the nucleotide sequence and
(b) detecting whether the probe or primer has specifically hybridized to the target polynucleotide, whereby specific hybridization provides a detection of the target polynucleotide in the sample.

25 19. The method of claim 16 or 18 wherein the polynucleotide probe or primer is a peptide nucleic acid.

20 20. A purified, recombinant Repro-EN-1.0 polypeptide whose amino acid sequence is identical to that of SEQ ID NO: 2, or an allelic variant of SEQ ID NO: 2.

25 21. A purified, recombinant IB1 polypeptide whose amino acid sequence is identical to that of SEQ ID NO: 4, or an allelic variant of SEQ ID NO: 4.

30 22. A purified polypeptide comprising an epitope of at least 5 amino acids of Repro-EN-1.0 (SEQ ID NO: 2), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.

35 23. A purified polypeptide comprising an epitope of at least 5 amino acids of IB1 (SEQ ID NO: 4), wherein the epitope specifically binds to antibodies from subjects diagnosed with endometriosis.

40 24. A composition consisting essentially of an antibody that specifically binds to Repro-EN-1.0 polypeptide (SEQ ID NO: 2).

45 25. A composition consisting essentially of an antibody that specifically binds to IB1 polypeptide (SEQ ID NO: 4).

26. The composition of claim 24 or 25 wherein the antibody is monoclonal.

45 27. A method for detecting a Repro-EN-1.0 polypeptide in a sample, comprising the steps of:

- 50 (a) contacting the sample with an antibody that specifically binds to the Repro-EN-1.0 polypeptide and
(b) detecting specific binding between the antibody and Repro-EN-1.0 polypeptide,

55 whereby specific binding provides a detection of Repro-EN-1.0 polypeptide in the sample.

28. A method for detecting a IB1 polypeptide in a sample, comprising the steps of:

- 55 (a) contacting the sample with an antibody that specifically binds to the IB1 polypeptide,
(b) detecting specific binding between the antibody and IB1 polypeptide,

whereby specific binding provides a detection of IB1 polypeptide in the sample.

29. A method for diagnosing endometriosis in a subject comprising the steps of:

(a) detecting a test amount of an antibody that specifically binds to Repro-EN-1.0 polypeptide in a sample from the subject; and

5 (b) comparing the test amount with a normal range of the antibody in a control sample from a subject who does not suffer from endometriosis,

10 whereby a test amount above the normal range provides a positive indication in the diagnosis of endometriosis.

15 30. The method of claim 29 wherein the step of detecting comprises capturing the antibody from the sample with an immobilized Repro-EN-1.0 or a peptide comprising an epitope of Repro-EN-1.0 and detecting captured antibody.

20 31. The method of claim 29 wherein the step of detecting comprises capturing the antibody from the sample with an immobilized anti-immunoglobulin antibody and detecting captured antibody.

25 32. The method of claim 30 wherein the step of detecting captured antibody comprises contacting the captured antibody with a detectable antibody that specifically binds immunoglobulins and detecting binding between the captured antibody and the detectable antibody.

30 33. The method of claim 31 wherein the step of detecting captured antibody comprises contacting the captured antibody with Repro-EN-1.0 or a polypeptide comprising an epitope of Repro-EN-1.0 and detecting binding between the captured antibody and the Repro-EN-1.0 or polypeptide.

35 34. A method for diagnosing endometriosis in a subject comprising the steps of:

(a) detecting a test amount of an antibody that specifically binds to IB1 polypeptide in a sample from the subject; and

40 (b) comparing the test amount with a normal range of the antibody in a control sample from a subject who does not suffer from endometriosis,

45 whereby a test amount above the normal range provides a positive indication in the diagnosis of endometriosis.

50 35. The method of any one of claims 29 to 34 wherein the sample comprises blood serum.

36. The method of any one of claims 29 to 34 wherein the antibody is an IgE immunoglobulin.

45 37. The method of any one of claims 29 to 34 wherein the antibody is an IgG immunoglobulin.

40 38. The method of any one of claims 29 to 34 wherein the antibody is an IgG₄ immunoglobulin.

45 39. The method of claim 34 wherein the step of detecting comprises capturing the antibody from the sample with an immobilized IB1 or a peptide comprising an epitope of IB1 and detecting captured antibody.

50 40. The method of claim 34 wherein the step of detecting comprises capturing the antibody from the sample with an immobilized anti-immunoglobulin antibody and detecting captured antibody.

55 41. The method of claim 39 wherein the step of detecting captured antibody comprises contacting the captured antibody with a detectable antibody that specifically binds immunoglobulins and detecting binding between the captured antibody and the detectable antibody.

42. The method of claim 40 wherein the step of detecting captured antibody comprises contacting the captured antibody with IB1 or a polypeptide comprising an epitope of IB1 and detecting binding between the captured antibody and the IB1 or polypeptide.

55 43. A method for use in following the progress of endometriosis in a subject comprising the steps of:

(a) detecting first and second amounts of an antibody that specifically binds Repro-EN-1.0 polypeptide in samples from the subject at a first and a second time, respectively; and
 (b) comparing the first and second amounts,

5 whereby an increase between the first and second amounts indicates progression of the endometriosis and a decrease between the first and second amounts indicates remission of the endometriosis.

44. A method for use in following the progress of endometriosis in a subject comprising the steps of:

10 (a) detecting first and second amounts of an antibody that specifically binds IB1 polypeptide in samples from the subject at a first and a second time, respectively; and
 (b) comparing the first and second amounts,

15 whereby an increase between the first and second amounts indicates progression of the endometriosis and a decrease between the first and second amounts indicates remission of the endometriosis.

45. An isolated MHC-peptide complex comprising:

20 at least a portion of an MHC Class I molecule or an MHC Class II molecule, wherein the portion comprises a binding site that specifically binds a peptide having an amino acid binding motif specific to the molecule, and wherein the portion engages in CD4-mediated or CD8-mediated binding to T cells, and
 a peptide of at least 8 amino acids in a sequence selected from the amino acid sequence of Repro-EN-1.0 (SEQ ID NO: 2), wherein the peptide comprises the amino acid binding motif and comprises an epitope that specifically binds to a T cell receptor;

25 wherein the complex specifically binds a T cell having a T cell receptor that specifically binds to the epitope, and wherein specific binding induces anergy in the T cell.

46. An isolated MHC-peptide complex comprising:

30 at least a portion of an MHC Class I molecule or an MHC Class II molecule, wherein the portion comprises a binding site that specifically binds a peptide having an amino acid binding motif specific to the molecule, and wherein the portion engages in CD4-mediated or CD8-mediated binding to T cells, and
 a peptide of at least 8 amino acids in a sequence selected from the amino acid sequence of IB 1 (SEQ ID NO: 4), wherein the peptide comprises the amino acid binding motif and comprises an epitope that specifically binds to a T cell receptor;

40 wherein the complex specifically binds a T cell having a T cell receptor that specifically binds to the epitope, and wherein specific binding induces anergy in the T cell.

47. An inhibitor of an immune response against Repro-EN-1.0 for use in a method for treating endometriosis in a subject.

48. The inhibitor of claim 47 which is an isolated MHC-peptide complex of claim 45 in an amount effective to inhibit the immune response.

49. The inhibitor of claim 47 which is an anti-idiotypic antibody that specifically binds to an antigen binding site of an antibody that specifically binds to Repro-EN-1.0 in an amount effective to inhibit the immune response.

50. An inhibitor of an immune response against IB1 for use in a method for treating endometriosis in a subject.

51. The inhibitor of claim 47 or 50 which is an immunosuppressant in an amount effective to inhibit the immune response.

52. The inhibitor of claim 50 which is an isolated MHC-peptide complex of claim 46 in an amount effective to inhibit the immune response.

53. The inhibitor of claim 50 which is an anti-idiotypic antibody that specifically binds to an antigen binding site of an

antibody that specifically binds to IB 1 in an amount effective to inhibit the immune response.

54. Use of an inhibitor of an immune response against Repro-EN-1.0 or IB1 for the manufacture of a medicament for treating endometriosis in a subject.

5 55. A screening method for determining whether a compound increases or decreases the expression of Repro-EN-1.0 in a cell, comprising contacting the cell with the compound and determining whether the production of Repro-EN-1.0 mRNA or polypeptide is increased or decreased.

10 56. A screening method for determining whether a compound increases or decreases the expression of IB1 in a cell, comprising contacting the cell with the compound and determining whether the production of IB1 mRNA or polypeptide is increased or decreased.

15 57. A method of detecting a chromosomal translocation of a Repro-EN-1.0 gene comprising the steps of:

15 (a) hybridizing a labelled polynucleotide probe that specifically hybridizes with the Repro-EN-1.0 nucleotide sequence of SEQ ID NO: 1 or its complement, to a chromosome spread from a cell sample to determine the pattern of hybridization and
20 (b) determining whether the pattern of hybridization differs from a normal pattern;

20 whereby a difference in the pattern represents a translocation.

58. A method of detecting a chromosomal translocation of a IB1 gene comprising the steps of:

25 (a) hybridizing a labelled polynucleotide probe that specifically hybridizes with the IB1 nucleotide sequence of SEQ ID NO: 3 or its complement, to a chromosome spread from a cell sample to determine the pattern of hybridization and
30 (b) determining whether the pattern of hybridization differs from a normal pattern;

30 whereby a difference in the pattern represents a translocation.

59. A method of detecting polymorphic forms of Repro-EN-1.0 comprising the steps of:

35 (a) determining the identity of a nucleotide or amino acid at a selected position within the sequence of a test Repro-EN-1.0 gene or polypeptide;
(b) determining the identity of the nucleotide or amino acid at the corresponding position of native Repro-EN-1.0 (SEQ ID NO: 1 or 2) gene or polypeptide; and
(c) comparing the identity from the test gene or polynucleotide with the identity of the native gene or polypeptide,
40 whereby a difference in identity indicates that the test polynucleotide is a polymorphic form of Repro-EN-1.0.

60. A method of detecting a polymorphic forms of IB1 comprising the steps of:

45 (a) determining the identity of a nucleotide or amino acid at a selected position within the sequence of a test IB1 gene or polypeptide;
(b) determining the identity of the nucleotide or amino acid at the corresponding position of native IB1 (SEQ ID NO: 3 or 4) gene or polypeptide; and
(c) comparing the identity from the test gene or polynucleotide with the identity of the native gene or polypeptide,
50 whereby a difference in identity indicates that the test polynucleotide is a polymorphic form of IB1.

FIG. 1.

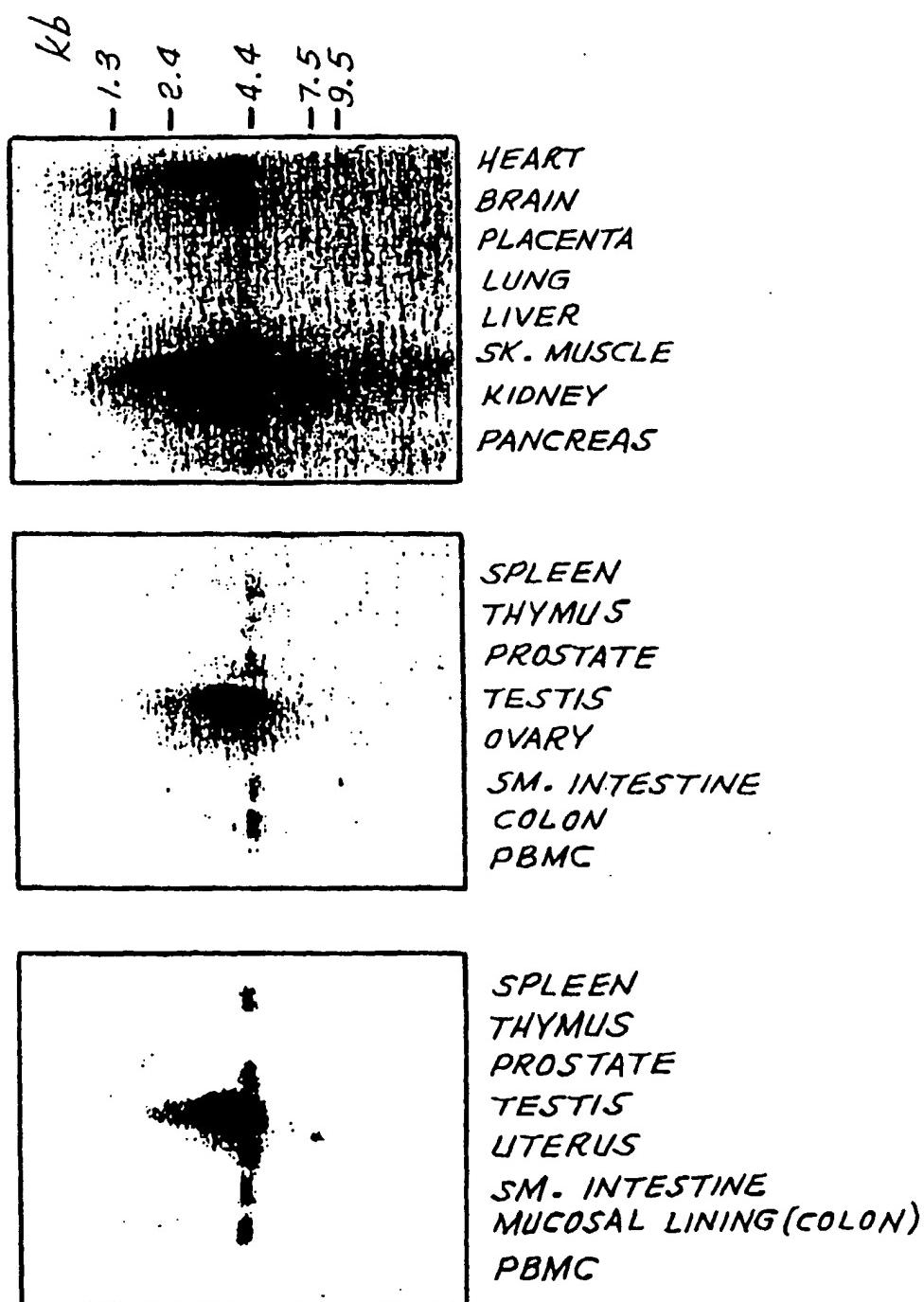


FIG. 2.

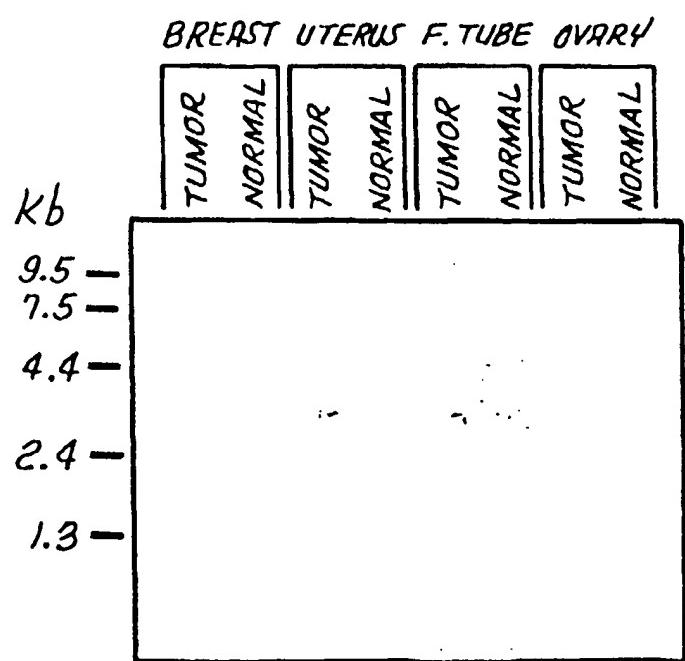
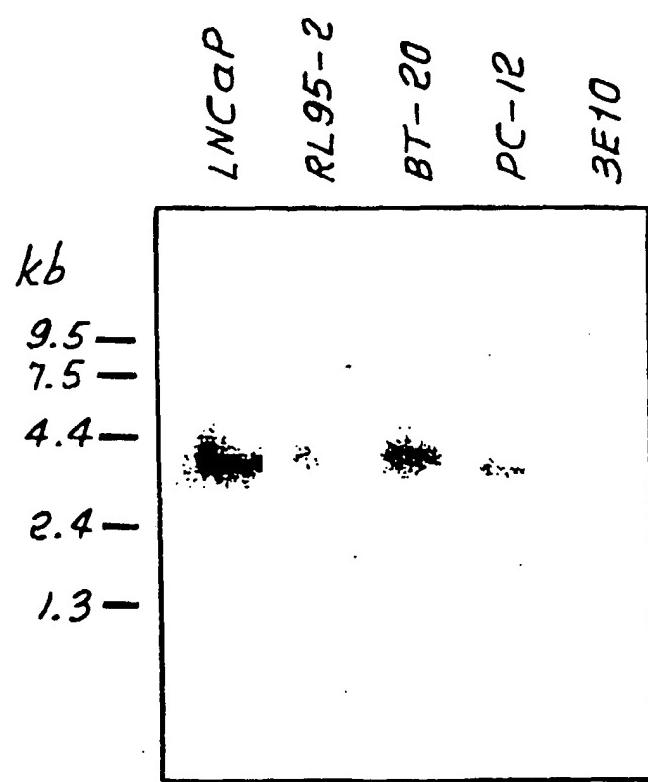


FIG. 3.



(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) EP 1 106 690 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
25.07.2001 Bulletin 2001/30

(51) Int Cl.7: C12N 15/12, C07K 14/47,
C12Q 1/68, C12N 5/10,
C07K 16/18, G01N 33/53,
G01N 33/557, C07K 19/00,
A61K 38/17

(21) Application number: 00310408.0

(22) Date of filing: 23.11.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE TR
Designated Extension States:
AL LT LV MK RO SI

- Menon, Surendra Nath
Culver City, California 90232 (US)
- French, Cynthia K.
Irvine, California 92612 (US)

(30) Priority: 23.11.1999 US 447399

(74) Representative:
Campbell, Patrick John Henry et al
J.A. Kemp & Co.,
14 South Square,
Gray's Inn
London WC1R 5JJ (GB)

(71) Applicant: DIAGNOSTIC PRODUCTS
CORPORATION
Los Angeles California 90045 (US)

(72) Inventors:
• El Shami, A. Said
Camarillo, California 93032 (US)

(54) Polynucleotide encoding autoantigens associated with endometriosis

(57) This invention provides a polynucleotide encoding Repro-EN-1.0 and IB1, polypeptides associated with endometriosis. Auto-antibodies against Repro-EN-

1.0 and IB1 have been found in subjects diagnosed with endometriosis. This invention also provides methods of using this polynucleotide and polypeptide.



European Patent
Office

PARTIAL EUROPEAN SEARCH REPORT

Application Number

which under Rule 45 of the European Patent Convention
shall be considered, for the purposes of subsequent
proceedings, as the European search report

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	Emest24 Database Entry Hs888330 Accession number W19888; 5 May 1996 HILLIER ET AL.: "The WashU-Merck EST Project" XP002126944 * the whole document * ---	16, 18, 57-60	C12N15/12 C07K14/47 C12Q1/68 C12N5/10 C07K16/18 G01N33/53 G01N33/557
X	EMBL Database Entry AI832097 Accession number AI832097; 13 July 1999 ROBERT STRAUSBERG XP002141979 * the whole document * ---	16, 18, 57-60	C07K19/00 A61K38/17
X	US 5 880 261 A (GERARD WAEBER ET AL.) 9 March 1999 (1999-03-09) * column 2, line 3 - column 4, line 27 * ---	28, 56	
A	WO 94 28021 A (MEDICAL UNIVERSITY OF SOUTH CAROLINA) 8 December 1994 (1994-12-08) * the whole document * -----	1-46, 48, 49, 52, 53, 55-60	
TECHNICAL FIELDS SEARCHED (Int.Cl.7)			
C12N C07K C12Q G01N A61K			
INCOMPLETE SEARCH			
<p>The Search Division considers that the present application, or one or more of its claims, does/do not comply with the EPC to such an extent that a meaningful search into the state of the art cannot be carried out, or can only be carried out partially, for these claims.</p> <p>Claims searched completely :</p> <p>Claims searched incompletely</p> <p>Claims not searched :</p> <p>Reason for the limitation of the search:</p> <p>see sheet C</p>			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	30 May 2001	Montero Lopez, B	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
<small>EPO FORM 1603.03.02 (EPC007)</small>			

European Patent
OfficeINCOMPLETE SEARCH
SHEET C

Application Number

EP 00 31 0408

Claim(s) searched completely:
1-46, 48, 49, 52, 53, 55-60

Claim(s) not searched:
47, 50, 51, 54

Reason for the limitation of the search:

Present claims 47, 50, 51 and 54 relate to a compound defined by reference to a desirable characteristic or property, namely inhibiting the immune response against Repro-EN-1.0 or IB1. The claims cover all compounds having this characteristic or property, whereas the application provides support within the meaning of Article 84 EPC and/or disclosure within the meaning of Article 83 EPC for only a very limited number of such compounds. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lack clarity (Article 84 EPC). An attempt is made to define the compound by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, no search has been carried out for claims 47, 50, 51 and 54.

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 31 0408

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

30-05-2001

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
US 5880261	A	09-03-1999	AU EP WO	6847298 A 0973895 A 9844106 A	22-10-1998 26-01-2000 08-10-1998
WO 9428021	A	08-12-1994	AU	6960694 A	20-12-1994